AD_____

Award Number: DAMD17-01-1-0736

TITLE: Plasticity of GABAergic Synapses in the Ventral Tegmental Area During Withdrawal From In Vivo Ethanol Administration

PRINCIPAL INVESTIGATOR: Antonello Bonci, M.D.

CONTRACTING ORGANIZATION: Ernest Gallo Clinic and Research Center
Emeryville, California 94608

REPORT DATE: July 2002

TYPE OF REPORT: Annual

PREPARED FOR: U.S. Army Medical Research and Materiel Command
Fort Detrick, Maryland 21702-5012

DISTRIBUTION STATEMENT: Approved for Public Release;
Distribution Unlimited

The views, opinions and/or findings contained in this report are those of the author(s) and should not be construed as an official Department of the Army position, policy or decision unless so designated by other documentation.

20021105 033

REPORT DOCUMENTATION PAGE

Form Approved
OMB No. 074-0188

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing this collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503

1. AGENCY USE ONLY (Leave blank)	2. REPORT DATE	3. REPORT TYPE AND DATES COVERED
	July 2002	Annual (1 Jul 01 - 30 Jun 02)

4. TITLE AND SUBTITLE
Plasticity of GABAergic Synapses in the Ventral Tegmental Area During Withdrawal From In Vivo Ethanol Administration

5. FUNDING NUMBERS
DAMD17-01-1-0736

6. AUTHOR(S)
Antonello Bonci, M.D.

7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES)
Ernest Gallo Clinic and Research Center
Emeryville, California 94608

E-Mail: bonci@itsa.ucsf.edu

8. PERFORMING ORGANIZATION REPORT NUMBER

9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES)
U.S. Army Medical Research and Materiel Command
Fort Detrick, Maryland 21702-5012

10. SPONSORING / MONITORING AGENCY REPORT NUMBER

11. SUPPLEMENTARY NOTES

12a. DISTRIBUTION / AVAILABILITY STATEMENT
Approved for Public Release; Distribution Unlimited

12b. DISTRIBUTION CODE

13. Abstract ***(Maximum 200 Words)*** *(abstract should contain no proprietary or confidential information)*
Ethanol consummatory behavior seems modulated through $GABA_A$ receptors in the ventral tegmental area (VTA). Previous exposure to ethanol enhances ethanol self-administration, but the mechanisms underlying this phenomenon is poorly understood. The purpose of Specific aim 1 was to examine changes occurring at GABA synapses onto VTA DA neurons after a single *in vivo* exposure to ethanol, and we have accomplished it. We observed that a single *in vivo* exposure to ethanol both shifted paired-pulse facilitation (PPF) to paired pulse depression (PPD) and increased ethanol self-administration. An increased frequency of spontaneous miniature $GABA_A$ IPSCs (mIPSCs) was also observed in the ethanol-treated animals. Further, the shift to PPD in the ethanol treated mice depended upon activation of presynaptic $GABA_B$ receptors. Forskolin increased the amplitude of $GABA_A$ IPSCs and the frequency of mIPSCs in the saline- but not in the ethanol-treated mice. Conversely, a protein kinase A (PKA) inhibitor significantly decreased both frequency of spontaneous mIPSCs and amplitude of $GABA_A$ IPSCs in the ethanol-treated mice. Our results indicate that potentiation of GABAergic synapses, via a PKA-dependent mechanism, occurs in the VTA after a single *in vivo* ethanol exposure, and such a potentiation might be a key synaptic modification underlying increased ethanol intake.

14. SUBJECT TERMS	15. NUMBER OF PAGES
GABA, VTA, ethanol, mesolimbic system, electrophysiology	24
	16. PRICE CODE

17. SECURITY CLASSIFICATION OF REPORT	18. SECURITY CLASSIFICATION OF THIS PAGE	19. SECURITY CLASSIFICATION OF ABSTRACT	20. LIMITATION OF ABSTRACT
Unclassified	Unclassified	Unclassified	Unlimited

Table of Contents

Introduction

The mesolimbic dopamine (DA) system, originating in the ventral tegmental area (VTA), is considered to be a central modulator of self-administration of all drugs of abuse, including ethanol. Specifically, we are interested in understanding the physiology of the $GABA_A$ synapses because the modulation of the $GABA_A$ receptor in the VTA plays a role in many ethanol-related behaviors, including ethanol self-administration. Our hypothesis is that a long-term enhancement of GABAergic synaptic transmission onto DA cells might be a central factor mediating the decrease in DA cell activity during withdrawal from ethanol; to prove it, we plan to characterize the short- and long-term changes of activity at $GABA_A$ receptors onto VTA DA cells after *in vivo* exposure to ethanol. We believe that an important part of reaching the ultimate goal of reducing ethanol consumption will be realized through a better characterization of the physiology of $GABA_A$ receptors in the VTA. A better knowledge of the intrinsic synaptic physiology of VTA GABAergic synapses will enable us to find novel targets aimed at counteracting long-term changes at inhibitory synapses caused by ethanol intake.

Research accomplishments:

This report refers to our first year of funding. We are confident that we have accomplished the goal of specific aim 1. The results of our experiments have been published very recently in a full-length paper in the *Journal of Neuroscience*. I have included a copy of our paper as an appendix, and will refer to it for the figures and for detailed descriptions of our results and their significance.

In specific aim 1, we proposed to characterize whether changes in GABAergic synaptic transmission occurred as a result of exposure to a single *in vivo* injection of ethanol. We proposed to perform these experiments both during acute (1-2 days) and sub-acute (5-30 days) withdrawal from ethanol.

First, we sought to determine whether changes in the probability of GABA release occurred at $GABA_A$ synapses in ethanol-treated mice. In order to accomplish this, we used the paired-pulse stimulation protocol to test for changes in synaptic strength elicited by paired stimuli given at an interval of 50 msec. As explained in our grant proposal, changes in transmitter release would generally affect the paired-pulse ratio (Khazipov et al., 1995, Mennerick and Zorumski, 1995; Debanne et al., 1996; Salin et al., 1996; Stoop and Poo, 1996; Murthy et al., 1997; Gottschalk et al., 1998; Hernandez-Echeagaray et al., 1998; Lessmann and Heumann, 1998; Emmerson and Miller, 1999; Niittykoski et al., 1999; Stanford and Cooper, 1999; Steffensen et al., 1999; Sullivan, 1999; Jiang et al., 2000; Poncer et al., 2000; Cooper and Stanford, 2001; Rozov et al., 2001). However, some inconsistencies have recently been reported for some brain areas (Brody and Yue, 2000; Kraushaar and Jonas, 2000; Waldeck et al., 2000). Figure 1 *a-e* of our paper (see appendix) shows paired-pulse facilitation (PPF) in the saline-treated mice, since the second pulse evoked a $GABA_A$ IPSC (IPSC2) that was significantly larger than the first ($IPSC2/IPSC1 = 1.3 \pm 0.06$; $n = 16$). Conversely, ethanol-treated mice exhibited paired

pulse depression (PPD; IPSC2/IPSC1 = 0.8 ± 0.02; $n = 17$; $p < 0.05$). Thus, after a single *in vivo* exposure to ethanol, the paired-pulse protocol resulted in PPD. Then, we decided to determine whether PPF or PPD were dependent on the initial size of the first $GABA_A$ IPSC. Our results show that PPF and PPD were not affected by changes in stimulus intensity (Fig. 1*d, appendix*). In addition, we analyzed the PPR by dividing the mean of IPSC2 by the mean of IPSC1, and as expected we observed similar results ($p < 0.05$; ethanol, IPSC2/IPSC1 = 0.7 ± 0.03; $n = 17$; saline, IPSC2/IPSC1 = 1.1 ± 0.06; $n = 16$).

Changes in probability of release, function of postsynaptic $GABA_A$ receptors, or a combination of these may be reflected by a variation in PPR toward PPD. We bath applied GABA (100 μM, 3 min) in the presence of the $GABA_B$ receptor antagonist (3-aminopropyl)(diethoxymethyl) phosphinic acid (CGP35348, 100 μM) to determine whether changes in $GABA_A$ receptor function, number, or both occurred following *in vivo* ethanol administration. The amplitude of the inward current elicited by GABA did not differ between ethanol- and saline-treated animals ($n = 4$ for each group; $p > 0.05$; Fig. 1*f*). These results suggest that modifications in transmitter release, rather than in postsynaptic receptors, occur after an *in vivo* exposure to ethanol. In the next series of experiments, we examined spontaneous $GABA_A$ mIPSCs. Figure 2 *a-c (see appendix)*, shows that the frequency of mIPSCs was significantly higher in ethanol- than in saline-treated animals (ethanol, 2.8 ± 0.4 Hz; $n = 9$; saline, 0.7 ± 0.1 Hz; $n = 7$; $p < 0.05$). Furthermore, there was no significant difference in the amplitude of mIPSCs in the two groups (mean amplitudes 30.4 ± 4.2 and 29.2 ± 3.4 pA in saline- and ethanol-treated mice, respectively Fig. 2 *c,d*; appendix, $p > 0.05$). To briefly summarize the experiments described so far, both the paired-pulse protocol and the increased frequency of spontaneous events indicate that the probability of GABA release in the VTA was increased 1 day after a single *in vivo* exposure to ethanol. Previous *in vivo* studies have shown that a decrease in DAergic activity lasts long after somatic signs of withdrawal have disappeared (Diana et al., 1996). Therefore, in order to determine how long the PPD lasted, we collected evoked $GABA_A$ IPSCs by using the paired-pulse protocol both 1 and 2 weeks after the single *in vivo* exposure to ethanol. Figure 1*e* (see appendix) shows that PPD was still present after 1 week (ethanol, IPSC2/IPSC1 = 0.9 ± 0.08; $n = 9$; $p < 0.05$; saline, IPSC2/IPSC1 = 1.2 ± 0.08; $n = 7$), but that within 2 weeks, the phenomenon had subsided (ethanol, IPSC2/IPSC1 = 1.1 ± 0.06; $n = 8$; $p > 0.05$; saline, IPSC2/IPSC1 = 1.2 ± 0.05; $n = 7$). These data are consistent with the idea that increased probability of GABA release might contribute to the long-lasting hypoactivity of DA neurons that has been observed *in vivo* after ethanol exposure (Diana et al., 1996).

Note: in specific aim 1, we proposed to study changes in GABAergic synaptic transmission both during acute (1-2 days) and sub-acute (5-30 days) withdrawal from ethanol. We did not perform any ratio measurement after day 15 since we did not detect any significant change in the paired-pulse ratio after 2 weeks from the ethanol injection.

In the next series of experiments, we proceeded to investigate in detail whether activation of presynaptic $GABA_B$ receptors was responsible for the observed difference in PPR between the two groups of animals. In the midbrain, $GABA_B$ receptors are present presynaptically and postsynaptically, and it has been shown that the activation of presynaptic $GABA_B$ receptors causes inhibition of $GABA_A$ IPSCs (Johnson and North, 1992b; Hausser and Yung, 1994). Therefore, we measured the PPR in the presence of the

$GABA_B$ receptor antagonist CGP35348 (100 μM, 5 min). Figure 4, *a* and *b (see appendix)*, shows that CGP35348 significantly shifted the PPD to PPF in ethanol-treated animals (IPSC2/IPSC1 = 0.7 ± 0.04-1.2 ± 0.08; $n = 10$; $p < 0.05$) by increasing the amplitude of the second evoked $GABA_A$ IPSC (IPSC2, 140 ± 9%; data not shown) without affecting either $GABA_A$ IPSC in the saline-treated animals (IPSC2/IPSC1 = 1.5 ± 0.1-1.4 ± 0.2; $n = 5$; $p > 0.05$).

In both groups of mice, the frequency and the amplitude of mIPSCs were unaffected by CGP35348 (100 μM, 5 min) (frequency: ethanol, 2.9 ± 0.2-2.6 ± 0.3 Hz; $n = 7$; saline, 1.1 ± 0.1-1.1 ± 0.1 Hz; $n = 7$; amplitude: ethanol, 33.6 ± 1.8-33 ± 2.9 pA; $n = 5$; saline, 33.2 ± 2.1-26.9 ± 2.6 pA; $n = 7$; Fig. 4*c,d*). In conclusion, the PPD observed in the ethanol-treated mice could result from an increased probability of GABA release in the ethanol- treated mice, and this might in turn lead to activation of presynaptic $GABA_B$ receptors. It is also possible that the sensitivity of presynaptic $GABA_B$ receptors might be enhanced in the ethanol-treated animals. In order to find out, we compared the inhibition caused by the $GABA_B$ receptor agonist baclofen in slices from both saline- and ethanol-treated animals by testing the differences in sensitivity of presynaptic $GABA_B$ receptors. The results shown in figure *5 a and b* (see appendix) show that the concentration-response curves to baclofen were similar in the saline- and ethanol-treated mice (baclofen 0.1 μM: ethanol, 22.3 ± 8.9%; $n = 5$; saline, 25.4 ± 3.1%; $n = 5$; baclofen 1 μM: ethanol, 42.8 ± 12.3%; $n = 5$; saline, 44.5 ± 2.3%; $n = 5$; baclofen 10 μM: ethanol, 70.6 ± 11.8%; $n = 5$; saline, 72.7 ± 7.2%; $n = 5$. Thus, the increase in the IPSC2 observed in the presence of CGP35348 in ethanol-treated animals did not result from altered sensitivity of $GABA_B$ receptors to endogenous GABA. Finally, when we applied a high dose of baclofen (10 μM; Fig. 5*c,d, see appendix*), such a dose reverted the PPD to PPF in ethanol-treated mice (IPSC2/IPSC1 = 0.7 ± 0.02-1.2 ± 0.1; $n = 5$; $p < 0.05$), but it produced an insignificant increase in PPF in saline-treated animals (IPSC2/IPSC1 = 1.2 ± 0.1-1.3 ± 0.1; $n = 5$). These results further support the hypothesis that GABA levels are increased after ethanol exposure, leading to spillover onto presynaptic $GABA_B$ receptors, whose activation leads to inhibition of GABA release.

Note: Because changes in the paired-pulse ratio and changes in the probability of release are generally linked, we expected baclofen to increase PPF in the saline group. However, we detected a non-significant increase in the PPF during the application of baclofen at this concentration (IPSC2/IPSC1 = 1.2 ± 0.09-1.3 ± 0.02; $n = 5$; $p > 0.05$) and at 1 μM (IPSC2/IPSC1 = 1.1 ± 0.07-1.2 ± 0.08; $n = 5$; $p > 0.05$) in the saline-treated animals. This indicates that the PPR measure is not sensitive enough in this range to detect a decrease in GABA release, possibly as a result of a ceiling effect.

In their 1997 work, Bonci and Williams described an increase in probability of GABA release during acute withdrawal from chronic morphine in several brain regions, including the VTA, as well as its dependence upon cAMP in the VTA. We decided to examine the possibility that the cAMP-dependent pathway was modified in the ethanol-treated animals since ethanol and other drugs of abuse are known to modulate the camp-protein kinase A (PKA) cascade within the mesolimbic system (Hoffman and Tabakoff, 1990; Self et al., 1998; Spanagel and Weiss, 1999). To do so, we directly activated adenylyl cyclase (AC) by bath applying forskolin. Forskolin (10 μM, 10 min) augmented

the IPSC1 in saline-treated animals (Fig. 6*a, see appendix,* 111.6 ± 17.4%; $n = 5$; $p < 0.05$) but had no effect on the amplitude of the IPSC1 in ethanol-treated mice (23.1 ± 13.3%; $n = 5$). Thus, application of forskolin decreased the paired-pulse ratio toward depression in slices from saline-treated animals (Fig. 6*b*; IPSC2/IPSC1 = 1.4 ± 0.1-0.9 ± 0.1; $n = 5$; $p < 0.05$), but was without effect in slices from ethanol-treated animals (IPSC2/IPSC1 = 0.8 ± 0.05-0.7 ± 0.1; $n = 5$). This supports the idea that activation of AC increased the probability of GABA release. In addition, the frequency of spontaneous mIPSCs was also significantly increased by forskolin in saline-treated mice (0.7 ± 0.1-1.7 ± 0.3 Hz; $n = 7$; $p < 0.05$; Fig. 6*c*), but not in the ethanol-treated animals (2.6 ± 0.7-2.9 ± 1.0 Hz; $n = 8$; Fig. 6*c*). There was no significant difference in the amplitude of the mIPSCs in the absence or presence of forskolin in slices from either group of animals (saline, 29.4 ± 2.5-31 ± 2.9 pA; $n = 7$; ethanol, 34 ± 4.2-26.5 ± 1.7 pA; $n = 8$; Fig. 6*d*). To rule out the possibility of nonspecific effects of forskolin, we tested 1,9-dideoxyforskolin, an inactive analog of forskolin (Seamon and Daly, 1985). Superfusion of 1,9-dideoxyforskolin (10 μM, 10 min) had no effect on IPSCs in either group of animals (7.7 ± 3.3 and 1.1 ± 2.6% in ethanol- and saline-treated mice, respectively; $n = 4$ for each group; data not shown). Taken together, these results suggest that a saturation of AC might occur after a single *in vivo* exposure to ethanol. Because PKA activity is altered by intracellular cAMP levels, in the next series of experiments we tested the effect of *N*-[2(*p*-bromocinnamylamino)ethyl]-5-isoquinolinesulfonamide (H89), which inhibits PKA in a competitive manner against ATP (Chijiwa et al., 1990). Superfusion of H89 (10 μM, 20 min) significantly reduced the amplitude of IPSC1 in ethanol-treated mice (38.6 ± 5.4%; $n = 5$; $p < 0.05$; Fig. 7*a, see appendix*), but had no effect in slices from control animals (3.4 ± 8.1%; $n = 5$). The subsequent application of forskolin (10 μM, 10 min) in the presence of H89, used to test the activity of H89 in control animals, did not change the amplitude of the IPSC1 in either group (Fig. 7*a, see appendix*). Furthermore, H89, by reducing the size of IPSC1, shifted the PPD to PPF in slices from ethanol-treated animals (IPSC2/IPSC1 = 0.8 ± 0.02-1.1 ± 0.01; $n = 5$; $p < 0.05$; Fig. 7*b, see appendix*). Consistent with these results, H89 reduced the frequency of spontaneous mIPSC in the ethanol-treated animals (2.6 ± 0.1-1.6 ± 0.1 Hz; $n = 6$; $p < 0.05$; Fig. 7*c, see appendix*), but not in the saline-treated animals (0.9 ± 0.1-0.8 ± 0.1 Hz; $n = 5$); in the presence of H89, the amplitude of spontaneous mIPSCs was not changed in either saline-treated mice (31.7 ± 5.2-31.3 ± 2.6 pA; $n = 5$) or ethanol-treated mice (28.7 ± 3.1-24.8 ± 2.9 pA; $n = 6$). These experiments indicate that a single exposure to ethanol enhances PKA activity, and that such a phenomenon increases the probability of GABA release in the VTA. In our final series of experiments, we decided to test whether changes in ethanol self-administration were produced by the same single *in vivo* ethanol exposure used for our electrophysiological experiments. Behavioral studies reported that a single *in vivo* exposure to ethanol produced an increase in subsequent self-administration of ethanol in rodents (Camarini et al., 2000; Files et al., 2000). Additionally, because $GABA_A$ agonists facilitate acquisition of voluntary ethanol drinking in rats (Smith et al., 1992; Nowak et al., 1998), activation of $GABA_A$ receptors plays a role in ethanol self-administration. Therefore, mice that underwent the same experimental protocol were studied for ethanol consumption. To perform these experiments, 24 hr after a single exposure to ethanol (2 gm/kg, i.p.), mice were given a two-bottle choice test with two concentrations of ethanol versus water for 5 d at each

concentration (2 and 5% v/v, respectively). In order to allow the animals to acclimate to the taste of ethanol, we began with the lower concentration for 5 d. As reported previously (Camarini et al., 2000), C57BL/6J mice pretreated with ethanol consumed more ethanol than saline controls after 10 d of housing in the continuous access situation (Fig. 3*a, see appendix*). We found that 10 d after the acute exposure, the mean ethanol intake was 9.3 ± 0.6 and 7.4 ± 0.4 gm/kg for ethanol-treated ($n = 8$) and saline-treated ($n = 8$) mice, respectively ($p < 0.05$). Although both groups preferred water to ethanol, ethanol-treated mice showed an increased preference for ethanol (46.9 ± 0.5%; $n = 8$; $p < 0.05$; Fig. 3*b, see appendix*) when compared with saline controls (41.1 ± 1.5%; $n = 8$). However, because increased ethanol intake observed in ethanol-treated mice may be affected by differential absorption, distribution, or clearance of ethanol, we measured blood ethanol concentrations 10-90 min after administration of ethanol (4 gm/kg, i.p.). Figure 3*c* of the appendix shows that ethanol clearance did not differ between the two groups of animals, although it showed lower blood ethanol levels than commonly reported (Hodge et al., 1999; Thiele et al., 2000; Wand et al., 2001). One possible explanation for such low levels might be the faster metabolism of younger mice used for the present study compared with the 2- to 4-month-old mice tested in other studies (Hodge et al., 1999; Thiele et al., 2000; Wand et al., 2001). In conclusion, taken together, our results support the hypothesis that an increased GABAergic transmission in the VTA may be involved in facilitating or maintaining ethanol consumption.

Key research accomplishment:

1) Our data demonstrate that a single *in vivo* injection of ethanol produces a long-term potentiation of GABAergic synapses onto dopaminergic neurons in the VTA.

2) Such a potentiation lasts for a week after the ethanol injection, and is caused by a cAMP-PKA dependent mechanism.

3) We have also provided evidence that a single *in vivo* injection of ethanol also increases subsequent ethanol consumption.

Conclusions:

To our knowledge, this is the first study that directly correlates a synaptic modification at $GABA_A$ receptors with an abnormal ethanol-dependent behavior, namely ethanol self-administration.

Our results suggest that increased $GABA_A$ receptor activation in the VTA plays a central role in increasing ethanol self-administration, and indicate that this cAMP-PKA-dependent

plasticity occurring at $GABA_A$ synapses might represent an important crucial signaling event underlying increased ethanol consumption.

Reportable outcomes:

One Full-length article:

1) "Long-lasting potentiation of GABAergic synapses in dopamine neurons after a single in vivo ethanol exposure". *J.Neuroscience*, 2002, 22 (6) 2074-82.

Abstracts:

1) Effects of ethanol withdrawal on gabaergic synapses in dopamine neurons. M. Melis, M.A. Ungless, & A. Bonci. Society for Neuroscience, Annual Conference, San Diego, 2001.

2) Effects of a single exposure to ethanol in mice: a behavioral and electrophysiological study. R. Camarini, M. Melis, C.W. Hodge & A. Bonci. Society for Neuroscience, Annual Conference, San Diego, 2001.

3) M. Melis, M.A. Ungless, A. Bonci (2001) The effects of alcohol withdrawal on synaptic transmission of VTA DA neurons in C57Bl/6J mice. 24th Annual Meeting of Research Society on Alcoholism, Montreal, June 23-28 2001.

Presentations:

Invited speaker in the seminar series at UT Southwestern, Dallas, TX (February 2002)
Invited speaker at the Annual Meeting of Research Society for Alcoholism (June 2002)
Invited speaker in the seminar series at UCSD, The Scripps Institute, San Diego (October 2002)
Invited speaker at Society for Neuroscience Symposium (November 2002)
Invited speaker at American College of Neuropsychopharmacology Symposium (December 2002).

REFERENCES

Bonci A, Williams JT (1997) Increased probability of GABA release during withdrawal from morphine. J Neurosci. 17(2): 796-803.

Brody DL, Yue DT (2000) Release-independent short-term synaptic depression in cultured hippocampal neurons. J Neurosci. 20(7):2480-94.

Camarini R, Mehmert KK , Hodge CW (2000) Effects of ethanol pre-exposure on ethanol self-administration in DBA/2J and C57BL/6 mice. Alcoholism Clinical and Experimental Research 24 (5) Suppl: 16A

Chijiwa T, Mishima A, Hagiwara M, Sano M, Hayashi K, Inoue T, Naito K, Toshioka T, Hidaka H (1990) Inhibition of forskolin-induced neurite outgrowth and protein phosphorylation by a newly synthesized selective inhibitor of cyclic AMP-dependent protein kinase, N-[2-(p-bromocinnamylamino)ethyl]-5-isoquinolinesulfonamide (H-89), of PC12D pheochromocytoma cells. J Biol Chem. 265(9): 5267-5272.

Cooper AJ, Stanford IM (2001) Dopamine D2 receptor mediated presynaptic inhibition of striatopallidal GABA(A) IPSCs in vitro. Neuropharmacology 41(1):62-71.

Debanne D, Guerineau NC, Gahwiler BH, Thompson SM (1996) Paired-pulse facilitation and depression at unitary synapses in rat hippocampus: quantal fluctuation affects subsequent release. J Physiol. 491 (Pt 1):163-176.

Diana M, Pistis M, Muntoni AL, Gessa GL (1996) Mesolimbic dopaminergic reduction outlasts ethanol withdrawal syndrome: evidence of protacted abstinence. Neuroscience 71(2): 411-415.

Emmerson PJ, Miller RJ (1999) Pre- and postsynaptic actions of opioid and orphan opioid agonists in the rat arcuate nucleus and ventromedial hypothalamus in vitro. J Physiol. 517 (Pt 2):431-45.

Files FJ, Samson HH, Denning CE (2000) Effects of prior ethanol exposure on ethanol self-administration in a continuous access situation using retractable drinking tubes. Alcohol 21(1): 97-102.

Gottschalk W, Pozzo-Miller LD, Figurov A, Lu B (1998) Presynaptic modulation of synaptic transmission and plasticity by brain-derived neurotrophic factor in the developing hippocampus. J Neurosci. 18(17):6830-9.

Hausser MA, Yung WH (1994) Inhibitory synaptic potentials in guinea-pig substantia nigra dopamine neurons in vitro. J Physiol 479 (3): 401-422.

Hernandez-Echeagaray E, Galarraga E, Bargas J (1998) 3-Alpha-chloro-imperialine, a potent blocker of cholinergic presynaptic modulation of glutamatergic afferents in the rat neostriatum. Neuropharmacology. 37(12):1493-502.

Hodge CW, Mehmert KK, Kelley SP, McMahon T, Haywood A, Olive MF, Wang D, Sanchez-Perez AM, Messing RO (1999) Supersensitivity to allosteric GABAA receptor modulators and alcohol in mice lacking PKCϵ. Nat Neurosci 2(11): 997-1002.

Hoffman PL,Tabakoff B (1990) Ethanol and guanine nucleotide binding proteins: a selective interaction. FASEB J 4:2612-2622.

Jiang L, Sun S, Nedergaard M, Kang J (2000) Paired-pulse modulation at individual GABAergic synapses in rat hippocampus. J Physiol. 523 Pt 2:425-39.

Johnson SW and North RA (1992b) Opioids excite dopamine neurons by hyperpolarization of local interneurons. J Neurosci 12: 483-488

Khazipov R, Congar P, Ben-Ari Y (1995) Hippocampal CA1 lacunosum-moleculare interneurons: modulation of monosynaptic GABAergic IPSCs by presynaptic GABAB receptors. J Neurophysiol. 74(5):2126-37.

Kraushaar U, Jonas P (2000) Efficacy and stability of quantal GABA release at a hippocampal interneuron-principal neuron synapse. J Neurosci. 20(15):5594-607.

Lessmann V, Heumann R (1998) Modulation of unitary glutamatergic synapses by neurotrophin-4/5 or brain-derived neurotrophic factor in hippocampal microcultures: presynaptic enhancement depends on pre-established paired-pulse facilitation. Neuroscience. 86(2):399-413.

Mennerick S, Zorumski CF (1995) Paired-pulse modulation of fast excitatory synaptic currents in microcultures of rat hippocampal neurons. J Physiol. 488 (Pt 1): 85-101.

Murthy VN, Sejnowski TJ, Stevens CF (1997) Heterogeneous release properties of visualized individual hippocampal synapses. Neuron. 18(4):599-612.

Niittykoski M, Ruotsalainen S, Haapalinna A, Larson J, Sirvio J (1999) Activation of muscarinic M3-like receptors and beta-adrenoceptors, but not M2-like muscarinic receptors or alpha-adrenoceptors, directly modulates corticostriatal neurotransmission in vitro. Neuroscience. 90(1):95-105.

Nowak KL, McBride WJ, Lumeng L, Li TK, Murphy JM (1998) Blocking GABAA receptors in the anterior ventral tegmental area attenuates ethanol intake of the alcohol-preferring P rats. Psychopharmacology 139 (1-2): 108- 116.

Poncer JC, McKinney RA, Gahwiler BH, Thompson SM (2000) Differential control of GABA release at synapses from distinct interneurons in rat hippocampus. J Physiol. 528 Pt 1:123-30.

Rozov A, Burnashev N, Sakmann B, Neher E (2001) Transmitter release modulation by intracellular Ca2+ buffers in facilitating and depressing nerve terminals of pyramidal cells in layer 2/3 of the rat neocortex indicates a target cell-specific difference in presynaptic calcium dynamics. J Physiol. 531(Pt 3):807-26.

Salin PA, Malenka RC, Nicoll RA (1996) Cyclic AMP mediates a presynaptic form of LTP at cerebellar parallel fiber synapses. Neuron 16(4): 797-803.

Seamon KB, Daly JW (1985). High-affinity binding of forskolin to rat brain membranes. Advances in Cyclic Nucleotide and Protein Phosphorylation Research 19: 125-135

Self DW, Genova LM, Hope BT, Barnhart WJ, Spencer JJ, Nestler EJ (1998) Involvement of cAMP- dependent protein kinase in the nucleus accumbens of cocaine self-administration and relapse of cocaine-seeking behavior. J Neurosci 18: 1848- 1859.

Smith BR, Robidoux J, Amit Z (1992) GABAergic involvement in the acquisition of voluntary ethanol intake in laboratory rats. Alcohol Alcohol. 27(3): 227-31.

Spanagel R, Weiss F (1999) The dopamine hypothesis of reward: past and current status. Trends Neurosci 22: 521-527.

Stanford IM, Cooper AJ (1999) Presynaptic mu and delta opioid receptor modulation of GABAA IPSCs in the rat globus pallidus in vitro. J Neurosci. 19(12):4796-803.

Steffensen SC, Henriksen SJ, Wilson MC (1999) Transgenic rescue of SNAP-25 restores dopamine-modulated synaptic transmission in the coloboma mutant. Brain Res. 847(2):186-95.

Stoop R, Poo MM (1996) Synaptic modulation by neurotrophic factors: differential and synergistic effects of brain-derived neurotrophic factor and ciliary neurotrophic factor. J Neurosci. 16(10):3256-64.

Sullivan JM (1999) Mechanisms of cannabinoid-receptor-mediated inhibition of synaptic transmission in cultured hippocampal pyramidal neurons. J Neurophysiol. 82(3):1286-94.

Thiele TE, Willias B, Stadler J, Reynolds JG, Bernestein IL, McKnight GS (2000) High ethanol consumption and low sensitivity to ethanol-induced sedation in protein kinase A-mutant mice. J Neurosci. 20: RC75.

Waldeck RF, Pereda A, Faber DS (2000) Properties and plasticity of paired-pulse depression at a central synapse. J Neurosci. 20(14):5312-20.

Wand G, Levine M, Zweifel L, Schwindinger W, Abel T (2001) The cAMP-Protein Kinase A Signal Transduction Pathway Modulates Ethanol Consumption and Sedative Effects of Ethanol. J Neurosci. 21(14): 5297-5303.

The Journal of Neuroscience, March 15, 2002, 22(6):2074–2082

Long-Lasting Potentiation of GABAergic Synapses in Dopamine Neurons after a Single *In Vivo* Ethanol Exposure

Miriam Melis, Rosana Camarini, Mark A. Ungless, and Antonello Bonci

Ernest Gallo Clinic and Research Center, Department of Neurology, University of California, San Francisco, California 94110

The mesolimbic dopamine (DA) system originating in the ventral tegmental area (VTA) is involved in many drug-related behaviors, including ethanol self-administration. In particular, VTA activity regulating ethanol consummatory behavior appears to be modulated through $GABA_A$ receptors. Previous exposure to ethanol enhances ethanol self-administration, but the mechanisms underlying this phenomenon are not well understood. In this study, we examined changes occurring at GABA synapses onto VTA DA neurons after a single *in vivo* exposure to ethanol. We observed that evoked $GABA_A$ IPSCs in DA neurons of ethanol-treated animals exhibited paired-pulse depression (PPD) compared with saline-treated animals, which exhibited paired-pulse facilitation (PPF). Furthermore, PPD was still present 1 week after the single exposure to ethanol. An increase in frequency of spontaneous miniature $GABA_A$ IPSCs (mIPSCs) was also observed in the ethanol-treated animals. Additionally, the $GABA_B$ receptor antagonist (3-aminopropyl) (diethoxymethyl) phosphinic acid shifted PPD to PPF, indicating that presynaptic $GABA_B$ receptor activation, likely attributable to GABA spillover, might play a role in mediating PPD in the ethanol-treated mice. The activation of adenylyl cyclase by forskolin increased the amplitude of $GABA_A$ IPSCs and the frequency of mIPSCs in the saline- but not in the ethanol-treated animals. Conversely, the protein kinase A (PKA) inhibitor *N*-[z-(*p*-bromocinnamylamino)ethyl]-5-isoquinolinesulfonamide significantly decreased both the frequency of spontaneous mIPSCs and the amplitude of $GABA_A$ IPSCs in the ethanol-treated mice but not in the saline controls. The present results indicate that potentiation of GABAergic synapses, via a PKA-dependent mechanism, occurs in the VTA after a single *in vivo* exposure to ethanol, and such potentiation might be a key synaptic modification underlying increased ethanol intake.

Key words: ventral tegmental area; ethanol; probability of GABA release; presynaptic plasticity; cAMP; PKA

The mesolimbic dopamine (DA) system originates in the ventral tegmental area (VTA), and it projects to structures associated with the limbic system, primarily the nucleus accumbens (NAcc), the amygdala, the hippocampus, and the prefrontal cortex (Fuxe et al., 1974; Oades and Halliday, 1987). A growing body of evidence implicates the mesolimbic DA system in the regulation of ethanol self-administration (Rassnick et al., 1993a; Samson et al., 1993; Ng and George, 1994; Koob et al., 1998; McBride et al., 1999; Kaczmarek and Kiefer, 2000; Nowak et al., 2000). An important role of DA in ethanol reinforcement has been suggested by studies showing that DA receptor antagonists, injected systemically or directly into the terminal regions of the mesolimbic DA system, decrease lever pressing for ethanol (Samson et al., 1993; Ng and George, 1994). Furthermore, a variety of pharmacological manipulations within this pathway, affecting the activity of DA neurons, produced changes in ethanol consumption, suggesting that DA neuronal activity within the VTA may be important for maintaining ethanol consummatory behavior (Rassnick et al., 1993a; Ng and George, 1994; Kaczmarek and Kiefer, 2000; Nowak et al., 2000). In fact, a marked reduction of the spontaneous activity of mesolimbic DA neurons (Diana et al., 1993; Bailey et al., 1998), resulting in decreased extracellular DA levels in NAcc (Diana et al., 1993), has been observed during acute withdrawal from chronic ethanol. Moreover, the fact that ethanol intake in dependent rats greatly exceeds that of nondependent rats during acute withdrawal, and that increased self-administration restores DA levels to normal in NAcc, suggests that decreased DA levels may trigger ethanol-seeking behavior (Weiss et al., 1996). The above-mentioned studies indicate that changes in activity of VTA DA cells, correlated with extracellular DA levels in the NAcc, might regulate ethanol consumption (McBride et al., 1995; Weiss et al., 1996; Hodge et al., 1997; Ikemoto et al., 1997), and accordingly, DA antagonists impair alcohol self-administration. In the midbrain dopamine systems, GABAergic neurons exert an inhibitory control on DA neurons (Johnson and North, 1992b; Hausser and Yung, 1994; Paladini et al., 1999). Therefore, hyperactivity of VTA GABA cells observed during acute withdrawal from chronic ethanol (Gallegos et al., 1999) could account, at least in part, for the reduced DAergic activity. Because the hypofunction of the DAergic system outlasts the somatic signs of acute withdrawal (Diana et al., 1996), such an increase of GABAergic synaptic transmission might not only represent a functional correlate of acute withdrawal from ethanol but also play a role in both short- and long-term consequences produced by ethanol exposure.

Interestingly, the influence of the initial exposure to ethanol and the patterns of its subsequent consumption have been ob-

Received Nov. 6, 2001; revised Dec. 17, 2001; accepted Dec. 28, 2001.

This work was supported by funds provided by the State of California for medical research on alcohol and substance abuse through the University of California, San Francisco, and by the Department of the Army, award number DAMD17-01-10736, United States Army Medical Research Acquisition Activity (Fort Detrick, MD). The content of the information does not necessarily reflect the position or the policy of the Government, and no official endorsement should be inferred. We thank all members of the Bonci Lab for many helpful discussions and inputs. We also deeply thank the late Dr. Thomas Dunwiddie for his many helpful insights. He will be sorely missed in the future.

Correspondence should be addressed to Antonello Bonci at the above address. E-mail: bonci@itsa.ucsf.edu.

Copyright © 2002 Society for Neuroscience 0270-6474/02/222074-09$15.00/0

served in both humans and laboratory animals (Haertzen et al., 1983; Camarini et al., 2000; Files et al., 2000). Unfortunately, the relationship between the reinforcing quality of the first experience and subsequent habits of ethanol consumption is still unclear. Whether this change in behavior is attributable to a reduced sensitivity to the stimulant effects of ethanol (Phillips et al., 1995) or a blockade of the development of ethanol-induced conditioned taste aversion (Risinger and Cunningham, 1995) remains to be elucidated. Because both systemic and intra-VTA administration of $GABA_A$ receptor agonists facilitate, whereas antagonists decrease, the acquisition of voluntary ethanol drinking in rats (Smith et al., 1992; Nowak et al., 1998), the GABAergic transmission within the VTA might play an important role.

Although the acute effects of ethanol in the mesolimbic system have been studied extensively (Brodie et al., 1990, 1999; Nie et al., 1993, 1994; Brodie and Appel, 1998, 2000; Steffensen et al., 2000), there are no studies directly examining whether synaptic changes occur in the VTA after exposure to ethanol.

To address this issue, we studied $GABA_A$-mediated IPSCs in VTA DA neurons 24 hr after a single injection of either ethanol (2 gm/kg, i.p.) or saline.

MATERIALS AND METHODS

Slice preparation. The preparation of VTA slices was as described previously (Thomas et al., 2000). Briefly, C57BL/6J mice (21–35 d; Charles River, Hollister, CA) were anesthetized with halothane and killed. A block of tissue containing the midbrain was sliced in the horizontal plane (230 μm) with a vibratome (Leica, Nussloch, Germany) in ice-cold low-Ca^{2+} solution containing (in mM): 126 NaCl, 1.6 KCl, 1.2 NaH_2PO_4, 1.2 MgCl, 0.625 $CaCl_2$, 18 $NaHCO_3$, and 11 glucose. Slices (two per animal) were transferred in a holding chamber with a bicarbonate-buffered solution (32–34°C) saturated with 95%O_2 and 5%CO_2 containing (in mM): 126 NaCl, 1.6 KCl, 1.2 NaH_2PO_4, 1.2 $MgCl_2$, 2.4 $CaCl_2$, 18 $NaHCO_3$, and 11 glucose. Slices were allowed to recover for at least 1 hr before being placed in the recording chamber and superfused with the bicarbonate-buffered solution (32–34°C) saturated with 95%O_2 and 5%CO_2.

Whole-cell recording. Only one cell for each experimental procedure was recorded per mouse. Cells were visualized with an upright microscope with infrared illumination, and whole-cell voltage-clamp recordings were made by using an Axopatch 1D amplifier (Axon Instruments, Foster City, CA). All $GABA_A$ IPSC recordings were made with electrodes filled with an internal solution containing the following (in mM): 144 KCl, 1 $CaCl_2$, 3.45 K_4BAPTA, 10 HEPES, 2 Mg_2ATP, and 0.25 Mg_2GTP, pH 7.2–7.4. Experiments were begun only after series resistance had stabilized (typically 15–40 MΩ). Series and input resistance were monitored continuously on-line with a 4 mV depolarizing step (25 msec). Data were filtered at 2 kHz, digitized at 10 kHz, and collected on-line with acquisition software (Igor Pro, Lake Oswego, OR). Because of the composition of the internal solution, the $GABA_A$ IPSCs were inward at a membrane potential of −70 mV and were completely blocked by picrotoxin (100 μM). DA cells were identified by the presence of a large I_h current (Johnson and North, 1992a) that was assayed immediately after break-in, using a series of incremental 10 mV hyperpolarizing steps from a holding potential of −70 mV. A bipolar stainless steel stimulating electrode was placed 100 μm rostral to the recording electrode and was used to stimulate at a frequency of 0.1 Hz. Neurons were voltage-clamped at a membrane potential of −70 mV. All $GABA_A$ IPSCs were recorded in the presence of 2-amino-5-phosphonopentanoic acid (AP5; 100 μM), 6-cyano-2,3-dihydroxy-7-nitro-quinoxaline (10 μM), strychnine (1 μM), and eticlopride (100 nM) to block NMDA, AMPA, glycine, and dopamine D2-mediated synaptic currents, respectively. As described previously (Bonci and Williams, 1997), this solution had no effect on the holding current of the dopamine cells. The amplitudes of IPSCs were calculated by taking a 1 msec window around the peak of the IPSC and comparing this with the 5 msec window immediately before the stimulation artifact. Paired stimuli were given with an interstimulus interval of 50 msec, and the ratio between the second and the first IPSCs was calculated and averaged for a 10 min baseline. Drugs were applied in known concentrations to the superfusion medium. The spontaneous miniature IPSCs (mIPSC) were collected in the presence of lidocaine (500 μM) and analyzed (120 sweeps for each condition, 1 sec/sweep) using Mini Analysis program (Synaptosoft). To accurately determine the mIPSC amplitude, only mIPSCs that were >8 pA were accepted for analysis. The choice of this cutoff amplitude for acceptance of mIPSCs was made to obtain a high signal-to-noise ratio.

Alcohol self-administration. Sixteen C57BL/6J mice (Charles River Laboratories, Wilmington, MA) were housed individually in polycarbonate cages, with food and water available *ad libitum*, and habituated to their home cage for 1 week before the experiment. The colony room was maintained on a 12 hr light/dark cycle with lights on at 6 A.M. All experimental procedures were conducted under institutional and National Institutes of Health guidelines. Oral ethanol self-administration was examined using a two-bottle choice protocol (Phillips et al., 1998; Hodge et al., 1999). Mice were offered the choice between 2% (v/v) ethanol versus water for 5 d. The ethanol concentration was increased to 5%, and the ethanol consumption was measured for 5 more days. Fluid volumes consumed were recorded every day, and the bottle positions were alternated daily. Each day, the mice were weighed, and then the ethanol consumption was calculated as grams of ethanol per kilogram.

Blood alcohol determination. Blood tail collection of 3- to 4-week-old mice did not provide enough volume for measurement; therefore, blood ethanol concentration was measured by drawing a 40 μl blood sample from the trunk. Twenty-four hours after either ethanol (2 gm/kg, i.p.) or saline exposure, blood samples were collected at 10, 30, 60, and 90 min after an intraperitoneal 4 gm/kg injection of ethanol (four or five mice were used for each group and for every time point). Blood plasma was extracted with trichloroacetic acid, and plasma ethanol content was measured using a 332 alcohol diagnostic kit (Sigma, St. Louis, MO).

Results in the text and figures are presented as the mean ± SEM. Results between groups were compared using a *t* test, either paired or unpaired where appropriate; $p < 0.05$ was taken as indicating statistical significance.

RESULTS

In the present study, we investigated the properties of $GABA_A$ IPSCs recorded in VTA DA cells from mice that received a single *in vivo* injection of ethanol (2 gm/kg, i.p.) or saline the day before the recordings.

First, to determine whether changes in the probability of GABA release occurred at these synapses in ethanol-treated mice, we use the paired-pulse stimulation protocol to test for changes in synaptic strength elicited by paired stimuli given at an interval of 50 msec. It has been shown that changes in transmitter release would generally affect the paired pulse ratio (Khazipov et al., 1995, Mennerick and Zorumski, 1995; Debanne et al., 1996; Salin et al., 1996; Stoop and Poo, 1996; Murthy et al., 1997; Gottschalk et al., 1998; Hernandez-Echeagaray et al., 1998; Lessmann and Heumann, 1998; Emmerson and Miller, 1999; Niitty-koski et al., 1999; Stanford and Cooper, 1999; Steffensen et al., 1999; Sullivan, 1999; Jiang et al., 2000; Poncer et al., 2000; Yun et al., 2000; Cooper and Stanford, 2001; Rozov et al., 2001). Although some inconsistencies have recently been reported for some brain areas (Brody and Yue, 2000; Kraushaar and Jonas, 2000; Waldeck et al., 2000), it is well established that changes in transmitter release affect the paired-pulse ratio (PPR) in the VTA (Bonci and Williams, 1997; Manzoni and Williams, 1999). In slices from saline-treated mice, we observed paired-pulse facilitation (PPF), with the second pulse evoking a $GABA_A$ IPSC (IPSC2) that was significantly larger than the first (Fig. 1*a*,*e*; IPSC2/IPSC1 = 1.3 ± 0.06; $n = 16$). Conversely, ethanol-treated mice exhibited paired pulse depression (PPD; Fig. 1*b*,*e*; IPSC2/IPSC1 = 0.8 ± 0.02; $n = 17$; $p < 0.05$). Thus, after a single *in vivo* exposure to ethanol, the paired-pulse protocol resulted in PPD (Fig. 1*c–e*). PPF or PPD did not depend on the size of the first $GABA_A$ IPSC (IPSC1; Fig. 1*c*) and was not affected by changes in stimulus intensity (Fig. 1*d*). In addition, we analyzed the PPR by dividing the mean of IPSC2 by the mean of IPSC1, and we find

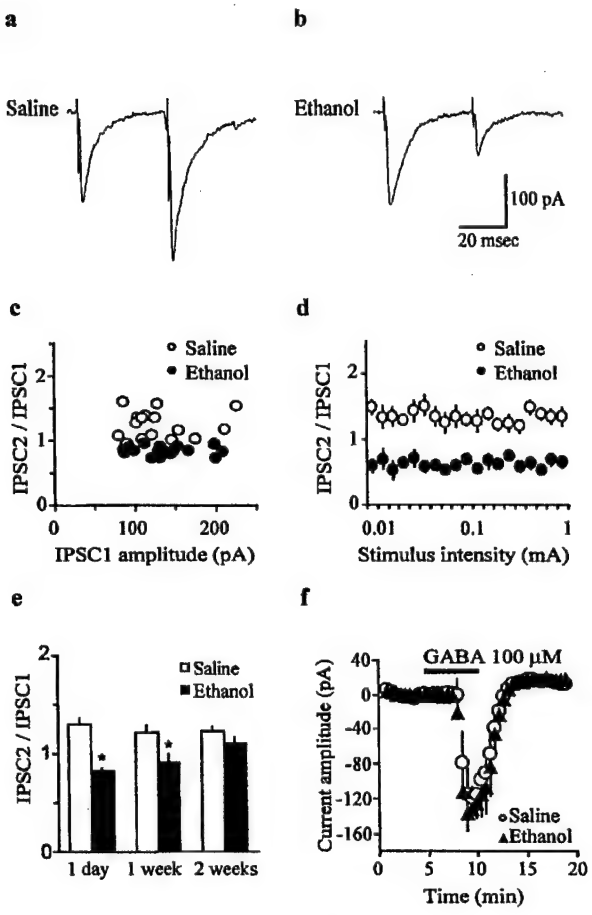

Figure 1. Increased probability of GABA release 24 hr after a single *in vivo* exposure to ethanol. $GABA_A$ IPSCs from ethanol-treated mice show PPD compared with saline controls, which show PPF. *a*, *b*, Examples of recordings from saline-treated (*a*) and ethanol-treated (*b*) animals. *c*, No correlation was found between the amplitude of IPSC1 and the IPSC2/IPSC1 ratio in both saline-treated mice ($n = 16$) and ethanol-treated mice ($n = 17$). *d*, The IPSC2/IPSC1 ratio is independent of the stimulus strength. Results are the average from four cells in each group of animals. *e*, PPD in ethanol-treated mice is a long-lasting phenomenon. The *bar graph* shows the average IPSC2/IPSC1 ratio (mean ± SEM) of saline- and ethanol-treated mice after 1 d ($n = 16$ and 17 for saline and ethanol, respectively; $^*p < 0.05$), 1 week ($n = 9$ per group; $^*p < 0.05$), and 2 weeks ($n = 8$ per group; $p > 0.05$) after ethanol pre-exposure. *f*, Ethanol pre-exposure does not change either number or function of postsynaptic $GABA_A$ receptors. Bath application of GABA (100 μM, 3 min) in the presence of $GABA_B$ receptor antagonist CGP 35348 (100 μM) elicited a similar current (30 sec bins) in both groups of animals ($n = 4$; $p > 0.05$) when neurons were voltage-clamped at −70 mV.

this method yields similar results ($p < 0.05$; ethanol, IPSC2/IPSC1 = 0.7 ± 0.03; $n = 17$; saline, IPSC2/IPSC1 = 1.1 ± 0.06; $n = 16$).

A change in PPR toward PPD might reflect changes in probability of release, function of postsynaptic $GABA_A$ receptors, or a combination of these. To determine whether changes in $GABA_A$ receptor function, number, or both occurred after *in vivo* ethanol administration, we bath applied GABA (100 μM, 3 min) in presence of the $GABA_B$ receptor antagonist (3-aminopropyl) (diethoxymethyl) phosphinic acid (CGP35348) (100 μM). The amplitude of the inward current elicited by GABA did not differ between ethanol- and saline-treated animals ($n = 4$ for each group; $p > 0.05$; Fig. 1*f*). These results suggest that modifications

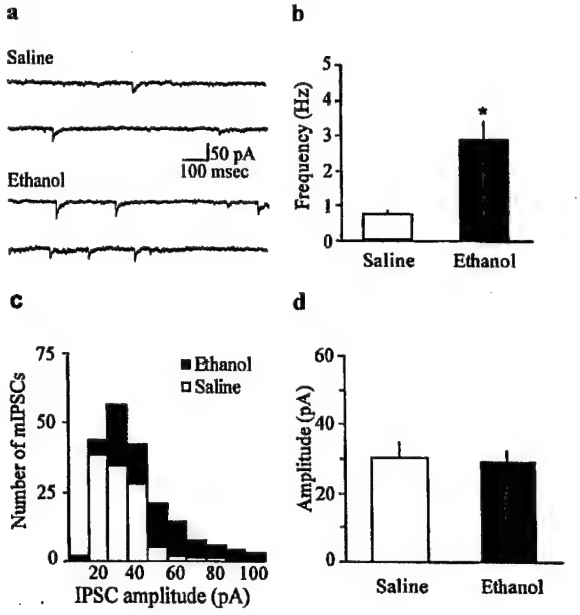

Figure 2. Ethanol pre-exposure increased the frequency, but not amplitude, of spontaneous mIPSCs. *a*, Samples of mIPSCs from saline-treated mice (*top traces*) and ethanol-treated mice (*bottom traces*). *b*, *Bar graph* showing the average (mean ± SEM) frequency for saline-treated animals ($n = 7$) and ethanol-treated animals ($n = 9$; $^*p < 0.05$). *c*, *Bar graph* (10 pA bins) showing an amplitude histogram of mIPSCs for ethanol-treated ($n = 9$) versus saline-treated ($n = 7$) mice. *d*, *Bar graph* showing the average (mean ± SEM) amplitude for saline-treated animals ($n = 7$) and ethanol-treated animals ($n = 9$).

in transmitter release, rather than in postsynaptic receptors, occur after an *in vivo* exposure to ethanol.

To further test this possibility, we examined spontaneous $GABA_A$ mIPSCs. Figure 2, *a–c*, shows that the frequency of mIPSCs was significantly higher in ethanol- than in saline-treated animals (ethanol, 2.8 ± 0.4 Hz; $n = 9$; saline, 0.7 ± 0.1 Hz; $n = 7$; $p < 0.05$). Furthermore, there was no significant difference in the amplitude of mIPSCs in the two groups, with mean amplitudes of 30.4 ± 4.2 and 29.2 ± 3.4 pA in saline- and ethanol-treated mice, respectively (Fig. 2*c*,*d*; $p > 0.05$). Because an increase in frequency but not amplitude of mIPSCs is generally thought to reflect a presynaptic increase in probability of transmitter release (Malenka and Nicoll, 1999), both the paired pulse protocol and the increased frequency of spontaneous events indicate that the probability of GABA release in the VTA was increased 1 d after a single exposure to ethanol.

Previous *in vivo* studies have shown that reduced DAergic activity persists long after somatic signs of withdrawal have subsided (Diana et al., 1996). Therefore, we collected evoked $GABA_A$ IPSCs by using the paired-pulse protocol both 1 and 2 weeks after the single *in vivo* exposure to ethanol to determine how long the PPD lasted. Figure 1*e* shows that PPD was still present after 1 week (ethanol, IPSC2/IPSC1 = 0.9 ± 0.08; $n = 9$; $p < 0.05$; saline, IPSC2/IPSC1 = 1.2 ± 0.08; $n = 7$), but that within 2 weeks, the phenomenon had subsided (ethanol, IPSC2/IPSC1 = 1.1 ± 0.06; $n = 8$; $p > 0.05$; saline, IPSC2/IPSC1 = 1.2 ± 0.05; $n = 7$). These data are consistent with the idea that increased probability of GABA release might contribute to the long-lasting hypoactivity of DA neurons that has been observed *in vivo* after ethanol exposure (Diana et al., 1996).

Behavioral studies reported that a single *in vivo* exposure to

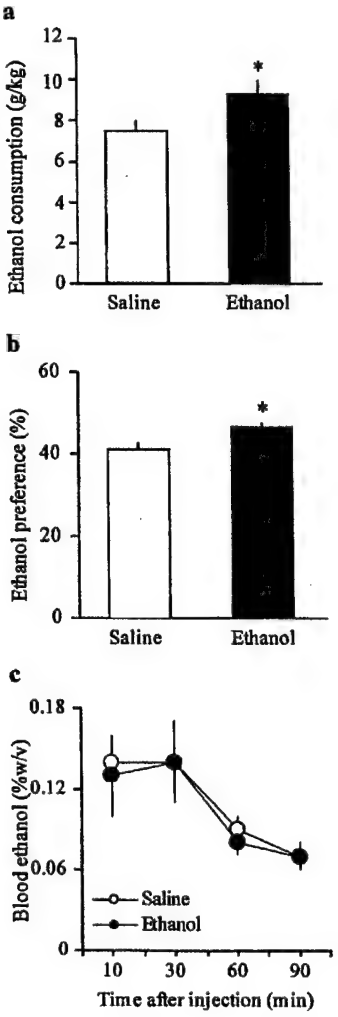

Figure 3. One week after the pre-exposure, ethanol-treated mice show increased ethanol intake and preference compared with saline controls. *a*, Voluntary 24 hr ethanol consumption (grams per kilogram) in C57BL/6J mice pretreated with ethanol ($n = 8$; $^*p < 0.05$) and saline ($n = 8$). *b*, Ethanol preference, calculated as 100 × milliliters of ethanol per total milliliters consumed. C57BL/6J mice pretreated with ethanol demonstrated a significant increase in ethanol preference ($n = 8$; $^*p < 0.05$) when compared with saline controls ($n = 8$). *c*, Blood ethanol clearance after acute administration of ethanol (4 gm/kg, i.p.) did not differ between ethanol- and saline-treated mice. Data (mean ± SEM) represent four animals per each group at every time point.

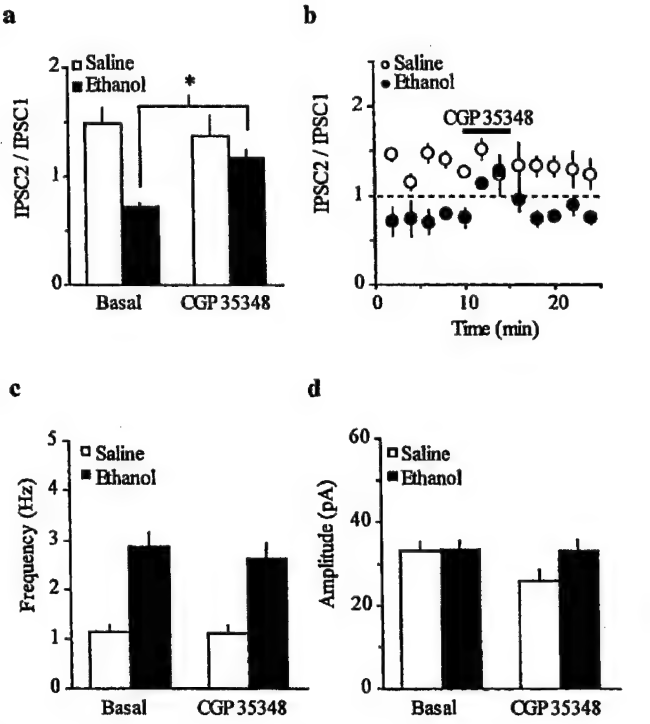

Figure 4. Effect of the $GABA_B$ receptor antagonist CGP35348 (100 μM) on IPSC2/IPSC1 ratio and spontaneous mIPSCs. *a*, CGP35348 (100 μM, 5 min) shifts the PPD to PPF in ethanol-treated mice ($n = 10$; $^*p < 0.05$), without affecting the PPF in saline-treated mice ($n = 5$). *b*, The IPSC2/IPSC1 ratio is plotted as a function of time in cells recorded from saline- and ethanol-treated mice and normalized against the mean of the first 10 min for each cell. *c*, CGP35348 does not change the frequency of mIPSCs in either group ($n = 7$ per each group). *d*, No changes in amplitude of spontaneous mIPSCs were found in either group ($n = 7$ per each group).

ethanol produced an increase in subsequent self-administration of ethanol in rodents (Camarini et al., 2000; Files et al., 2000). In addition, activation of $GABA_A$ receptors plays a role in ethanol self-administration, because $GABA_A$ agonists facilitate acquisition of voluntary ethanol drinking in rats (Smith et al., 1992; Nowak et al., 1998). We therefore decided to measure the ethanol consumption in mice that underwent the same experimental protocol. To perform these experiments, 24 hr after a single exposure to ethanol (2 gm/kg, i.p.), mice were given a two-bottle choice test with two concentrations of ethanol versus water for 5 d at each concentration (2 and 5% v/v, respectively). We began with the lower concentration for 5 d to allow the animals to acclimate to the taste of ethanol. As reported previously (Camarini et al., 2000), C57BL/6J mice pretreated with ethanol consumed more ethanol than saline controls after 10 d of housing in the continuous access situation (Fig. 3*a*). We found that 10 d after the acute exposure, the mean ethanol intake was 9.3 ± 0.6 and 7.4 ± 0.4 gm/kg for ethanol-treated ($n = 8$) and saline-treated ($n = 8$) mice, respectively ($p < 0.05$). Although both groups preferred water to ethanol, ethanol-treated mice showed an increased preference for ethanol (46.9 ± 0.5%; $n = 8$; $p < 0.05$; Fig. 3*b*) when compared with saline controls (41.1 ± 1.5%; $n = 8$). However, because differential absorption, distribution, or clearance of ethanol may contribute to the increased ethanol intake observed in ethanol-treated mice, we measured blood ethanol concentrations 10–90 min after administration of ethanol (4 gm/kg, i.p.). Figure 3*c* shows that ethanol clearance did not differ between the two groups of animals, although it showed lower blood ethanol levels than commonly reported (Hodge et al., 1999; Thiele et al., 2000; Wand et al., 2001). One possible explanation for such low levels might be the faster metabolism of younger mice used for the present study compared with the 2- to 4-month-old mice tested in other studies (Hodge et al., 1999; Thiele et al., 2000; Wand et al., 2001). Nevertheless, taken together, these results support the hypothesis that an increased GABAergic transmission in the VTA may be involved in facilitating or maintaining ethanol consumption (Smith et al., 1992; Nowak et al., 1998).

The observed difference in PPR between the two groups of animals might result indirectly from activation of presynaptic receptors. In the midbrain, $GABA_B$ receptors are present presynaptically and postsynaptically, and it has been shown that the activation of presynaptic $GABA_B$ receptors causes inhibition of

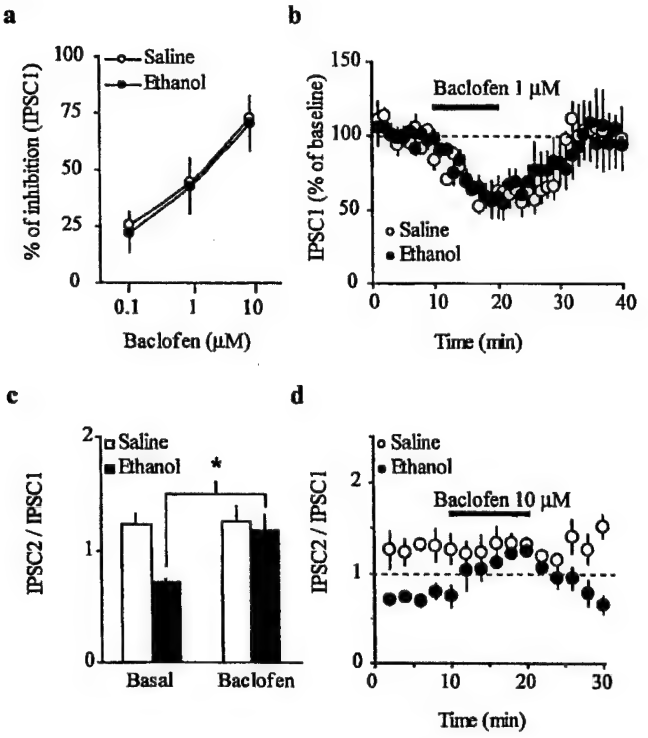

Figure 5. Effect of the $GABA_B$ receptor agonist baclofen on evoked IPSCs. *a*, Concentration–response curve for baclofen measuring the amplitude of IPSC1 from saline- and ethanol-treated animals ($n = 5$ per each group at all doses tested). *b*, Baclofen (1 μM, 10 min) decreases the amplitude of IPSC1 in both saline- and ethanol-treated mice to the same extent ($n = 5$ per each group). *c*, Baclofen (10 μM, 10 min) shifts the PPD to PPF in ethanol-treated mice ($n = 5$; $^*p < 0.05$) without affecting the PPF in saline-treated mice ($n = 5$). *d*, IPSC2/IPSC1 ratio plotted as a function of time in cells recorded from saline- and ethanol-treated mice ($n = 5$ per group; $^*p < 0.05$) and normalized against the mean of the first 10 min for each cell.

$GABA_A$ IPSCs (Johnson and North, 1992b; Hausser and Yung, 1994). Therefore, to test the possibility that increased probability of GABA release might raise GABA levels and thus activate presynaptic $GABA_B$ receptors, we measured the PPR in the presence of the $GABA_B$ receptor antagonist CGP35348 (100 μM, 5 min). Figure 4, *a* and *b*, shows that CGP35348 significantly shifted the PPD to PPF in ethanol-treated animals (IPSC2/IPSC1 = 0.7 ± 0.04–1.2 ± 0.08; $n = 10$; $p < 0.05$) by increasing the amplitude of the second evoked $GABA_A$ IPSC (IPSC2, 140 ± 9%; data not shown) without affecting either $GABA_A$ IPSC in the saline-treated animals (IPSC2/IPSC1 = 1.5 ± 0.1–1.4 ± 0.2; $n = 5$; $p > 0.05$). In addition, both the frequency and the amplitude of mIPSCs were unaffected by CGP35348 (100 μM, 5 min) in both groups of mice (frequency: ethanol, 2.9 ± 0.2–2.6 ± 0.3 Hz; $n = 7$; saline, 1.1 ± 0.1–1.1 ± 0.1 Hz; $n = 7$; amplitude: ethanol, 33.6 ± 1.8–33 ± 2.9 pA; $n = 5$; saline, 33.2 ± 2.1–26.9 ± 2.6 pA; $n = 7$; Fig. 4*c*,*d*). Thus, the PPD observed in the ethanol-treated mice could result from an increased probability of GABA release, which might in turn lead to activation of presynaptic $GABA_B$ receptors and decrease the IPSC2.

An alternative interpretation of the present results is that the sensitivity of presynaptic $GABA_B$ receptors might be enhanced in the ethanol-treated animals. Therefore, we tested differences in sensitivity of presynaptic $GABA_B$ receptors by comparing the inhibition caused by the $GABA_B$ receptor agonist baclofen in slices from both saline- and ethanol-treated animals. Figure 5*a* shows that the concentration–response curves to baclofen were similar in the saline- and ethanol-treated mice (baclofen 0.1 μM: ethanol, 22.3 ± 8.9%; $n = 5$; saline, 25.4 ± 3.1%; $n = 5$; baclofen 1 μM: ethanol, 42.8 ± 12.3%; $n = 5$; saline, 44.5 ± 2.3%; $n = 5$; baclofen 10 μM: ethanol, 70.6 ± 11.8%; $n = 5$; saline, 72.7 ± 7.2%; $n = 5$). Because the amplitude of the IPSC1 was decreased in both groups of animals to the same extent at all doses tested (Fig. 5*a*,*b*), we concluded that the increase in the IPSC2 observed in the presence of CGP35348 in ethanol-treated animals did not result from altered sensitivity of $GABA_B$ receptors to endogenous GABA. Additionally, a high dose of baclofen (10 μM; Fig. 5*c*,*d*) reverted the PPD to PPF in ethanol-treated mice (IPSC2/IPSC1 = 0.7 ± 0.02–1.2 ± 0.1; $n = 5$; $p < 0.05$), but it produced a nonsignificant increase in PPF in saline-treated animals (IPSC2/IPSC1 = 1.2 ± 0.1–1.3 ± 0.1; $n = 5$). These results further support the hypothesis that GABA levels are increased after ethanol exposure, leading to spillover onto presynaptic $GABA_B$ receptors, whose activation leads to inhibition of release (Hausser and Yung, 1994). Because changes in the paired-pulse ratio generally reflect changes in the probability of release, we expected baclofen to increase PPF in the saline group. However, we did detect a nonsignificant increase in the PPF during the application of baclofen at this concentration (IPSC2/IPSC1 = 1.2 ± 0.09–1.3 ± 0.02; $n = 5$; $p > 0.05$) and at 1 μM (IPSC2/IPSC1 = 1.1 ± 0.07–1.2 ± 0.08; $n = 5$; $p > 0.05$) in the saline-treated animals. This indicates that the PPR measure is not sensitive enough in this range to detect a decrease in GABA release, possibly as a result of a ceiling effect.

An increase in probability of GABA release has been described previously during acute withdrawal from chronic morphine in several brain regions, including the VTA (Bonci and Williams, 1997). In particular, this phenomenon has been characterized as being cAMP-dependent in the VTA (Bonci and Williams, 1997), the periaqueductal gray (Ingram et al., 1998), the NAcc (Chieng and Williams, 1998), and the dorsal raphe nucleus (Jolas et al., 2000). Ethanol and other drugs of abuse are known to modulate the cAMP–protein kinase A (PKA) cascade within the mesolimbic system (Hoffman and Tabakoff, 1990; Self et al., 1998; Spanagel and Weiss, 1999). Therefore, to examine the possibility that the cAMP-dependent pathway was modified in the ethanol-treated animals, we directly activated adenylyl cyclase (AC) by bath applying forskolin. Forskolin (10 μM, 10 min) augmented the IPSC1 in saline-treated animals (Fig. 6*a*; 111.6 ± 17.4%; $n = 5$; $p < 0.05$) but had no effect on the amplitude of the IPSC1 in ethanol-treated mice (23.1 ± 13.3%; $n = 5$). Thus, application of forskolin decreased the paired-pulse ratio toward depression in slices from saline-treated animals (Fig. 6*b*; IPSC2/IPSC1 = 1.4 ± 0.1–0.9 ± 0.1; $n = 5$; $p < 0.05$) but was without effect in slices from ethanol-treated animals (IPSC2/IPSC1 = 0.8 ± 0.05–0.7 ± 0.1; $n = 5$). This supports the idea that activation of AC increased the probability of GABA release. In addition, the frequency of spontaneous mIPSCs was also significantly increased by forskolin in saline-treated mice (0.7 ± 0.1–1.7 ± 0.3 Hz; $n = 7$; $p < 0.05$; Fig. 6*c*) but not in the ethanol-treated animals (2.6 ± 0.7–2.9 ± 1.0 Hz; $n = 8$; Fig. 6*c*). There was no significant difference in the amplitude of the mIPSCs in the absence or presence of forskolin in slices from either group of animals (saline, 29.4 ± 2.5–31 ± 2.9 pA; $n = 7$; ethanol, 34 ± 4.2–26.5 ± 1.7 pA; $n = 8$; Fig. 6*d*). To rule out the possibility of nonspecific effects of forskolin, we tested 1,9-dideoxyforskolin, an inactive analog of forskolin (Seamon and Daly, 1985). Superfusion of 1,9-dideoxyforskolin

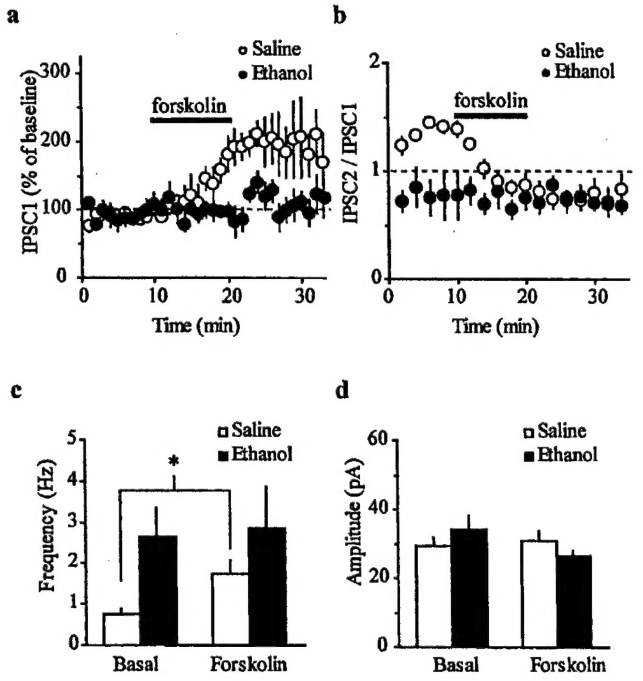

Figure 6. Effect of forskolin on evoked and spontaneous IPSCs. *a*, Forskolin (10 μM, 10 min) increases the amplitude of evoked IPSC1 in saline-treated mice ($n = 5$; $^*p < 0.05$) but not ethanol-treated mice ($n = 5$). *b*, Forskolin (10 μM, 10 min) shifts the PPF to PPD in saline-treated mice ($n = 5$; $^*p < 0.05$) without affecting the PPD in ethanol-treated mice ($n = 5$). The IPSC2/IPSC1 ratio is plotted as function of time in cells recorded from saline- and ethanol-treated mice and normalized against the mean of the first 10 min for each cell. *c*, Forskolin (10 μM, 10 min) induces a significant increase in the frequency of mIPSCs in saline-treated mice ($n = 7$; $^*p < 0.05$) but not in ethanol-treated mice ($n = 9$). *d*, No changes in amplitude were found in either group.

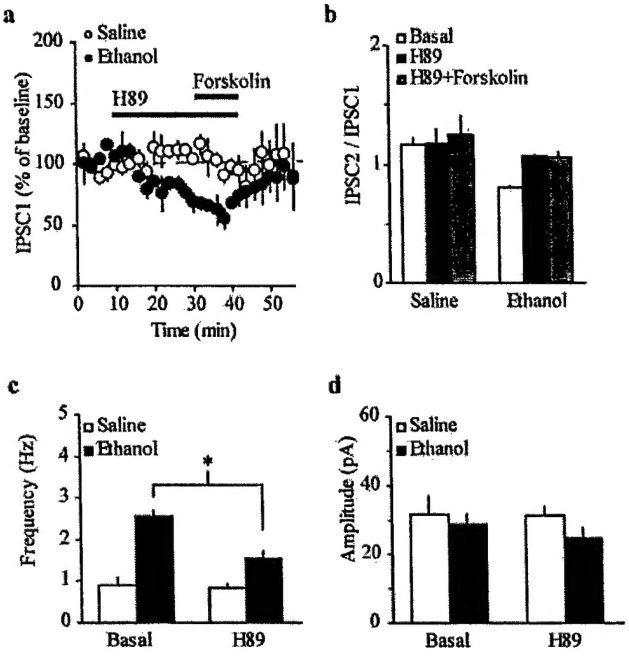

Figure 7. Effect of H89 on evoked and spontaneous IPSCs. *a*, H89 (10 μM, 20 min) decreases the amplitude of evoked IPSCs in ethanol-treated mice ($n = 5$; $^*p < 0.05$), but not saline-treated mice ($n = 5$). *b*, H89 (10 μM, 20 min) shifts the PPD to PPF in ethanol-treated mice ($n = 5$; $^*p < 0.05$) without affecting the PPF in saline-treated mice. *c*, H89 (10 μM, 20 min) induces a significant decrease in the frequency of mIPSCs in ethanol-treated mice ($n = 6$; $^*p < 0.05$) but not saline-treated mice ($n = 5$). *d*, No changes in amplitude were found in either group.

(10 μM, 10 min) had no effect on IPSCs in either group of animals (7.7 ± 3.3 and 1.1 ± 2.6% in ethanol- and saline-treated mice, respectively; $n = 4$ for each group; data not shown). Taken together, these results suggest that a saturation of AC might occur after a single *in vivo* exposure to ethanol.

Because changes of intracellular cAMP levels subsequently alter PKA activity, we tested the effect of *N*-[2(*p*-bromocinnamylamino)ethyl]-5-isoquinolinesulfonamide (H89), which inhibits PKA in a competitive manner against ATP (Chijiwa et al., 1990). Superfusion of H89 (10 μM, 20 min) significantly reduced the amplitude of IPSC1 in ethanol-treated mice (38.6 ± 5.4%; $n = 5$; $p < 0.05$; Fig. 7*a*) but had no effect in slices from control animals (3.4 ± 8.1%; $n = 5$). The subsequent application of forskolin (10 μM, 10 min) in the presence of H89, used to test the activity of H89 in control animals, did not change the amplitude of the IPSC1 in either group (Fig. 7*a*). Furthermore, H89, by reducing the size of IPSC1, shifted the PPD to PPF in slices from ethanol-treated animals (IPSC2/IPSC1 = 0.8 ± 0.02–1.1 ± 0.01; $n = 5$; $p < 0.05$; Fig. 7*b*). Consistent with these results, H89 reduced the frequency of spontaneous mIPSC in the ethanol-treated animals (2.6 ± 0.1–1.6 ± 0.1 Hz; $n = 6$; $p < 0.05$; Fig. 7*c*) but not in the saline-treated animals (0.9 ± 0.1–0.8 ± 0.1 Hz; $n = 5$); in the presence of H89, the amplitude of spontaneous mIPSCs was not changed in either saline-treated mice (31.7 ± 5.2–31.3 ± 2.6 pA; $n = 5$) or ethanol-treated mice (28.7 ± 3.1–24.8 ± 2.9 pA; $n = 6$). In conclusion, these experiments indicate that PKA activity is significantly enhanced by a single exposure to ethanol, and that such a phenomenon increases the probability of GABA release in the VTA.

DISCUSSION

In the present study, we observe that a single *in vivo* exposure to ethanol produces a long-lasting increase in the probability of GABA release in the VTA, and that such an increase is dependent on the activation of the cAMP–PKA signaling cascade. We hypothesize that this type of plasticity may play an important role in determining the increased alcohol consumption observed after a single exposure to ethanol (Spanagel and Weiss, 1999; Camarini et al., 2000). Our data, together with the fact that we and others have observed increased ethanol consumption when mice were pre-exposed to ethanol, indicate that increased probability of GABA release and increased ethanol self-administration, both produced by the single ethanol injection, might be strictly associated. Indeed, activation of $GABA_A$ receptors plays a role in ethanol self-administration, because $GABA_A$ agonists facilitate acquisition of voluntary ethanol drinking in rats (Smith et al., 1992; Nowak et al., 1998). Accordingly, a role of $GABA_A$ receptors within the VTA in mediating ethanol intake has been suggested (Samson et al., 1987; Boyle et al., 1993; Rassnick et al., 1993b). Indeed, systemic administration of $GABA_A$ receptor antagonists reduces intake (Boyle et al., 1993) and operant responding for ethanol in rats (Samson et al., 1987; Rassnick et al., 1993b). Consistent with these and previous findings, intra-VTA infusions of $GABA_A$ receptor antagonists decreased ethanol consumption in rats of the alcohol-preferring P line (Nowak et al., 1998). To further support the hypothesis that increased GABAergic activity produced by a single *in vivo* exposure to ethanol plays

a role in ethanol-related behaviors, it has recently been shown that there is a direct relationship between pretreatment with ethanol and enhanced self-administration of ethanol in mice (Camarini et al., 2000). Specifically, C57BL/6J mice pre-exposed to ethanol exhibited a significant increase of ethanol intake, and DBA/2J mice, which normally avoid oral ingestion of ethanol, did start to self-administer ethanol in a two-bottle choice test.

In our first set of experiments, we show that a single injection of ethanol shifted the paired-pulse modulation of $GABA_A$ IPSCs from PPF to PPD. The paired pulse stimulation is typically used as an electrophysiological protocol to test for changes in probability of transmitter release (Zucker, 1989; Stuart and Redman, 1991; Manabe et al., 1993; Mennerick and Zorumski, 1995; Debanne at al., 1996). Although this phenomenon is not always use-dependent (Brody and Yue, 2000; Kraushaar and Jonas, 2000; Waldeck et al., 2000), a variety of manipulations that increase transmitter release, including exposure to drugs of abuse, have been found to shift the paired-pulse ratio from facilitation toward depression in the hippocampus (Mennerick and Zorumski, 1995; Salin et al., 1996) and the VTA (Bonci and Williams, 1997). Furthermore, the persistence of PPD 1 week after the ethanol injection suggests that increased $GABA_A$-mediated inhibition may be considered a measure of changes occurring at these synapses, eventually contributing to the expression of ethanol-seeking behavior.

However, an increase in the probability of GABA release might simply be one of many factors determining the shift from PPF to PPD in the ethanol-treated animals. Although a desensitization of postsynaptic $GABA_A$ receptors could account for PPD, we tend to rule out that possibility, because the ethanol-treated animals show an increase in mISPCs frequency but not in amplitude when compared with the saline-treated animals. Furthermore, bath application of GABA, in the presence of a $GABA_B$ receptor antagonist, produces similar responses in saline- and ethanol-treated animals.

An alternative explanation for the observed PPD in the ethanol-treated animals, is that activation of presynaptic $GABA_B$ receptors might occur as a consequence of increased GABA levels produced by the first evoked stimulus, thus reducing the amplitude of IPSC2. Indeed, it has been shown that activation of $GABA_B$ receptors inhibits $GABA_A$ IPSCs in the midbrain via a presynaptic mechanism and therefore are considered to serve also as autoreceptors (Hausser and Yung, 1994). Our results showing that the $GABA_B$ antagonist CGP35348 shifted PPD to PPF in animals injected with ethanol, but not in the saline controls, indicate that presynaptic $GABA_B$ receptors might play a minor role when probability of GABA release is relatively low, as in the saline-injected animals. However, they might act as a negative feedback mechanism to regulate GABAergic transmission within the VTA when probability of GABA release is increased, such as after a single *in vivo* exposure to ethanol. Thus, our data indicate that increased GABA levels, by changing the spatial range of synaptically released GABA, allow the activation of presynaptic $GABA_B$ receptors located on the GABAergic interneurons, which in turn would prevent excessive $GABA_A$-mediated synaptic transmission (McCarren and Alger, 1985; Deisz and Prince, 1989; Davies et al., 1990; Isaacson et al., 1993).

Our results also suggest that the ethanol-induced increase in the probability of GABA release was a result of saturation of the AC cascade within GABAergic terminals. Forskolin, which enhanced the amplitude of evoked IPSCs and the frequency of mIPSCs in saline-treated mice, had no effect in the ethanol-treated mice. In addition, the PKA inhibitor H89 reduced the amplitude of evoked IPSCs and the frequency of mIPSCs only in ethanol-treated animals, whereas it had no effect in saline-treated animals. These findings suggest that a single *in vivo* exposure to ethanol results in persistent enhancement of PKA-dependent processes in GABAergic terminals in the VTA. It is possible that because of high cAMP levels after the exposure to ethanol, the catalytic subunits of the PKA complex become unbound and freely diffuse within the terminals. In fact, in slices from ethanol-treated animals, H89 revealed an increased basal activity of PKA, and the activation of AC by forskolin was blunted. Consistent with our findings, reduced signaling through the cAMP–PKA system, whether because of decreased expression of the α subunit of the stimulatory G-protein (Gs_α) or inhibition of PKA, changed C57BL/6J mice, considered to be an ethanol-preferring line of mice, into ethanol nonpreferring mice (Wand et al., 2001). In addition, alcohol-preferring rats show increased AC activity and expression of Gs_α in mesolimbic regions when compared with alcohol-nonpreferring rats (Froehlich and Wand, 1997). More generally, our results are in agreement with previous studies reporting that genetic manipulations of the cAMP–PKA pathway modulate ethanol intake and sensitivity to its sedative effects (Thiele et al., 2000; Wand et al., 2001). Although the relationships between the sedative and rewarding effects of ethanol are complex, it is also important to mention that the cAMP–PKA system has been implicated in neural plasticity associated with drug tolerance and dependence (Self and Nestler, 1995; Moore et al., 1998; Andretic et al., 1999; Yoshimura and Tabakoff, 1999). Finally, it has been shown that chronic exposure to many drugs of abuse, including ethanol, also leads to increased activity of cAMP-dependent processes (Terwilliger et al., 1991; Dohrman et al., 1996; Bonci and Williams, 1997).

In conclusion, our results provide evidence that a single *in vivo* exposure to ethanol produces a long-lasting potentiation of GABAergic synapses in the VTA. This cAMP–PKA-dependent plasticity occurring at these synapses might represent an important cellular signaling event underlying increased ethanol consumption. Whether these changes are the result of a compensation for acute effects of ethanol or manifestation of a long-lasting effect of acute ethanol remains to be elucidated. Although acute effects of ethanol on the GABAergic systems in the CNS are still a matter of debate, we tend to support the latter possibility, because ethanol has been found to enhance GABAergic transmission in several brain regions (Celentano et al., 1988; Deitrich et al., 1989; Aguayo and Pancetti, 1994; Mehta and Ticku, 1994; Wan et al., 1996; Nie et al., 2000), including the VTA (M. Melis and A. Bonci, unpublished observations). In conclusion, we observed that a single *in vivo* ethanol exposure induces a long-lasting potentiation of GABA synaptic transmission within the VTA via a cAMP–PKA mechanism, and that it appears to be directly related to increased ethanol consumption and may therefore be involved in the development of alcoholism.

REFERENCES

Aguayo LG, Pancetti FC (1994) Ethanol modulation of the gamma-aminobutyric acidA- and glycine-activated Cl− current in cultured mouse neurons. J Pharmacol Exp Ther 270:61–69.

Andretic R, Chaney S, Hirsh J (1999) Requirement of circadian genes for cocaine sensitization in *Drosophila*. Science 285:1066–1068.

Bailey CP, Manley SJ, Watson WP, Wonnacott S, Molleman A, Little HJ (1998) Chronic ethanol administration alters activity in ventral tegmental area neurons after cessation of withdrawal hyperexcitability. Brain Res 803:144–152.

Bonci A, Williams JT (1997) Increased probability of GABA release during withdrawal from morphine. J Neurosci 17:796–803.

Boyle AE, Segal R, Smith BR, Amit Z (1993) Bidirectional effects of GABAergic agonists and antagonists on maintenance of voluntary ethanol intake in laboratory rats. Pharmacol Biochem Behav 46:179–182.

Brodie MS, Appel SB (1998) The effects of ethanol on dopaminergic neurons of the ventral tegmental area studied with intracellular recording in brain slices. Alcohol Clin Exp Res 22:236–244.

Brodie MS, Appel SB (2000) Dopaminergic neurons in the ventral tegmental area of C57BL/6J and DBA/2J mice differ in sensitivity to ethanol excitation. Alcohol Clin Exp Res 24:1120–1124.

Brody DL, Yue DT (2000) Release-independent short-term synaptic depression in cultured hippocampal neurons. J Neurosci 20:2480–2494.

Brodie MS, Shefner SA, Dunwiddie TV (1990) Ethanol increases the firing rate of dopamine neurons of the rat ventral tegmental area in vitro. Brain Res 508:65–69.

Brodie MS, Pesold C, Appel SB (1999) Ethanol directly excites dopaminergic ventral tegmental area reward neurons. Alcohol Clin Exp Res 23:1848–1852.

Brundege JM, Dunwiddie TV (1996) Modulation of excitatory synaptic transmission by adenosine released from single hippocampal pyramidal neurons. J Neurosci 16:5603–5612.

Camarini R, Mehmert KK, Hodge CW (2000) Effects of ethanol preexposure on ethanol self-administration in DBA/2J and C57BL/6 mice. Alcohol Clin Exp Res [Suppl] 24:16A.

Celentano JJ, Gibbs TT, Farb DH (1988) Ethanol potentiates GABA- and glycine-induced chloride currents in chick spinal cord neurons. Brain Res 455:377–380.

Chieng B, Williams JT (1998) Increased opioid inhibition of GABA release in nucleus accumbens during morphine withdrawal. J Neurosci 18:7033–7039.

Chijiwa T, Mishima A, Hagiwara M, Sano M, Hayashi K, Inoue T, Naito K, Toshioka T, Hidaka H (1990) Inhibition of forskolin-induced neurite outgrowth and protein phosphorylation by a newly synthesized selective inhibitor of cyclic AMP-dependent protein kinase, *N*-[2-(*p*-bromocinnamylamino)ethyl]-5-isoquinolinesulfonamide (H-89), of PC12D pheochromocytoma cells. J Biol Chem 265:5267–5272.

Cooper AJ, Stanford IM (2001) Dopamine D2 receptor mediated presynaptic inhibition of striatopallidal GABA(A) IPSCs in vitro. Neuropharmacology 41:62–71.

Davies CH, Davies SN, Collingridge GL (1990) Paired-pulse depression of monosynaptic GABA-mediated inhibitory postsynaptic responses in rat hippocampus. J Physiol (Lond) 424:513–531.

Debanne D, Guerineau NC, Gahwiler BH, Thompson SM (1996) Paired-pulse facilitation and depression at unitary synapses in rat hippocampus: quantal fluctuation affects subsequent release. J Physiol (Lond) 491:163–176.

Deisz RA, Prince DA (1989) Frequency-dependent depression of inhibition in guinea-pig neocortex in vitro by GABAB receptor feed-back on GABA release. J Physiol (Lond) 412:513–541.

Deitrich RA, Dunwiddie TV, Harris RA, Erwin VG (1989) Mechanism of action of ethanol: initial central nervous system actions. Pharmacol Rev 41:489–537.

Diana M, Pistis M, Carboni S, Gessa GL, Rossetti ZL (1993) Profound decrement of mesolimbic dopaminergic neuronal activity during ethanol withdrawal syndrome in rats: electrophysiological and biochemical evidence. Proc Natl Acad Sci USA 90:7966–7969.

Diana M, Pistis M, Muntoni AL, Gessa GL (1996) Mesolimbic dopaminergic reduction outlasts ethanol withdrawal syndrome: evidence of protracted abstinence. Neuroscience 71:411–415.

Dohrman DP, Diamond I, Gordon AS (1996) Ethanol causes translocation of cAMP-dependent protein kinase catalytic subunit to the nucleus. Proc Natl Acad Sci USA 93:10217–10221.

Emmerson PJ, Miller RJ (1999) Pre- and postsynaptic actions of opioid and orphan opioid agonists in the rat arcuate nucleus and ventromedial hypothalamus in vitro. J Physiol (Lond) 517:431–445.

Files FJ, Samson HH, Denning CE (2000) Effects of prior ethanol exposure on ethanol self-administration in a continuous access situation using retractable drinking tubes. Alcohol 21:97–102.

Froehlich JC, Wand GS (1997) Adenylyl cyclase signal transduction and alcohol-induced sedation. Pharmacol Biochem Behav 58:1021–1030.

Fuxe K, Nystrom M, Tovi M, Smith R, Ogren SO (1974) Central catecholamine neurons, behavior and neuroleptic drugs: an analysis to understand the involvement of catecholamines in schizophrenia. J Psychiatr Res 11:151–161.

Gallegos RA, Lee RS, Criado JR, Henriksen SJ, Steffensen SC (1999) Adaptive responses of gamma-aminobutyric acid neurons in the ventral tegmental area to chronic ethanol. J Pharmacol Exp Ther 291:1045–1053.

Gottschalk W, Pozzo-Miller LD, Figurov A, Lu B (1998) Presynaptic modulation of synaptic transmission and plasticity by brain-derived neurotrophic factor in the developing hippocampus. J Neurosci 18:6830–6839.

Haertzen CA, Kocher TR, Miyasato K (1983) Reinforcements from the first drug experience can predict later drug habits and/or addiction: results with coffee, cigarettes, alcohol, barbiturates, minor and major tranquilizers, stimulants, marijuana, hallucinogens, heroin, opiates and cocaine. Drug Alcohol Depend 11:147–165.

Hausser MA, Yung WH (1994) Inhibitory synaptic potentials in guinea-pig substantia nigra dopamine neurons in vitro. J Physiol (Lond) 479:401–422.

Hernandez-Echeagaray E, Galarraga E, Bargas J (1998) 3-Alpha-chloroimperialine, a potent blocker of cholinergic presynaptic modulation of glutamatergic afferents in the rat neostriatum. Neuropharmacology 37:1493–1502.

Hodge CW, Samson HH, Chappelle AM (1997) Alcohol self-administration: further examination of the role of dopamine receptors in the nucleus accumbens. Alcohol Clin Exp Res 21:1083–1091.

Hodge CW, Mehmert KK, Kelley SP, McMahon T, Haywood A, Olive MF, Wang D, Sanchez-Perez AM, Messing RO (1999) Supersensitivity to allosteric $GABA_A$ receptor modulators and alcohol in mice lacking PKCε. Nat Neurosci 2:997–1002.

Hoffman PL, Tabakoff B (1990) Ethanol and guanine nucleotide binding proteins: a selective interaction. FASEB J 4:2612–2622.

Ikemoto S, McBride WJ, Murphy JM, Lumeng L, Li TK (1997) 6-OHDA-lesions of the nucleus accumbens disrupt the acquisition but not the maintenance of ethanol consumption in the alcohol-preferring P line of rats. Alcohol Clin Exp Res 21:1042–1046.

Ingram SL, Vaughan CW, Bagley EE, Connor M, Christie MJ (1998) Enhanced opioid efficacy in opioid dependence is caused by an altered signal transduction pathway. J Neurosci 18:10269–10276.

Isaacson JS, Solis JM, Nicoll RA (1993) Local and diffuse synaptic actions of GABA in the hippocampus. Neuron 10:165–175.

Jiang L, Sun S, Nedergaard M, Kang J (2000) Paired-pulse modulation at individual GABAergic synapses in rat hippocampus. J Physiol (Lond) 523:425–439.

Johnson SW, North RA (1992a) Two types of neurone in the rat ventral tegmental area and their synaptic inputs. J Physiol (Lond) 450:455–468.

Johnson SW, North RA (1992b) Opioids excite dopamine neurons by hyperpolarization of local interneurons. J Neurosci 12:483–488.

Jolas T, Nestler EJ, Aghajanian GK (2000) Chronic morphine increases GABA tone on serotonergic neurons of the dorsal raphe nucleus: association with an up-regulation of the cyclic AMP pathway. Neuroscience 95:433–443.

Kaczmarek HJ, Kiefer SW (2000) Microinjections of dopaminergic agents in the nucleus accumbens affect ethanol consumption but not palatability. Pharmacol Biochem Behav 66:307–312.

Khazipov R, Congar P, Ben-Ari Y (1995) Hippocampal CA1 lacunosum-moleculare interneurons: modulation of monosynaptic GABAergic IPSCs by presynaptic GABAB receptors. J Neurophysiol 74:2126–2137.

Koob GF, Roberts AJ, Schulteis G, Parsons LH, Heyser CJ, Hyytia P, Merlo-Pich E, Weiss F (1998) Neurocircuitry targets in ethanol reward and dependence. Alcohol Clin Exp Res 22:3–9.

Kraushaar U, Jonas P (2000) Efficacy and stability of quantal GABA release at a hippocampal interneuron-principal neuron synapse. J Neurosci 20:5594–5607.

Lessmann V, Heumann R (1998) Modulation of unitary glutamatergic synapses by neurotrophin-4/5 or brain-derived neurotrophic factor in hippocampal microcultures: presynaptic enhancement depends on preestablished paired-pulse facilitation. Neuroscience 86:399–413.

Malenka RC, Nicoll RA (1999) Long-term potentiation—a decade of progress? Science 285:1870–1874.

Manabe T, Wyllie DJ, Perkel DJ, Nicoll RA (1993) Modulation of synaptic transmission and long-term potentiation: effects on paired pulse facilitation and EPSC variance in the CA1 region of the hippocampus. J Neurophysiol 70:1451–1459.

Manzoni OJ, Williams JT (1999) Presynaptic regulation of glutamate release in the ventral tegmental area during morphine withdrawal. J Neurosci 19:6629–6636.

McBride WJ, Bodart B, Lumeng L, Li TK (1995) Association between low contents of dopamine and serotonin in the nucleus accumbens and high alcohol preference. Alcohol Clin Exp Res 19:1420–1422.

McBride WJ, Murphy JM, Ikemoto S (1999) Localization of brain reinforcement mechanisms: intracranial self-administration and intracranial place-conditioning studies. Behav Brain Res 101:129–152.

McCarren M, Alger BE (1985) Use-dependent depression of IPSPs in rat hippocampal pyramidal cells in vitro. J Neurophysiol 53:557–571.

Mehta AK, Ticku MK (1994) Ethanol enhancement of GABA-induced 36Cl− influx does not involve changes in Ca2+. Pharmacol Biochem Behav 47:355–357.

Mennerick S, Zorumski CF (1995) Paired-pulse modulation of fast excitatory synaptic currents in microcultures of rat hippocampal neurons. J Physiol (Lond) 488:85–101.

Moore MS, DeZazzo J, Luk AY, Tully T, Singh CM, Heberlein U (1998) Ethanol intoxication in *Drosophila*: genetic and pharmacological evidence for regulation by the cAMP signaling pathway. Cell 93:997–1007.

Murthy VN, Sejnowski TJ, Stevens CF (1997) Heterogeneous release properties of visualized individual hippocampal synapses. Neuron 18:599–612.

Ng GY, George SR (1994) Dopamine receptor agonist reduces ethanol self-administration in the ethanol-preferring C57BL/6J inbred mouse. Eur J Pharmacol 269:365–374.
Nie Z, Yuan X, Madamba SG, Siggins GR (1993) Ethanol decreases glutamatergic synaptic transmission in rat nucleus accumbens in vitro: naloxone reversal. J Pharmacol Exp Ther 266:1705–1712.
Nie Z, Madamba SG, Siggins GR (1994) Ethanol inhibits glutamatergic neurotransmission in nucleus accumbens neurons by multiple mechanisms. J Pharmacol Exp Ther 271:1566–1573.
Nie Z, Madamba SG, Siggins GR (2000) Ethanol enhances gamma-aminobutyric acid responses in a subpopulation of nucleus accumbens neurons: role of metabotropic glutamate receptors. J Pharmacol Exp Ther 293:654–661.
Niittykoski M, Ruotsalainen S, Haapalinna A, Larson J, Sirvio J (1999) Activation of muscarinic M3-like receptors and beta-adrenoceptors, but not M2-like muscarinic receptors or alpha-adrenoceptors, directly modulates corticostriatal neurotransmission in vitro. Neuroscience 90:95–105.
Nowak KL, McBride WJ, Lumeng L, Li TK, Murphy JM (1998) Blocking GABAA receptors in the anterior ventral tegmental area attenuates ethanol intake of the alcohol-preferring P rats. Psychopharmacology 139:108–116.
Nowak KL, McBride WJ, Lumeng L, Li TK, Murphy JM (2000) Involvement of dopamine D2 autoreceptors in the ventral tegmental area on alcohol and saccharin intake of the alcohol-preferring P rat. Alcohol Clin Exp Res 24:476–483.
Oades RD, Halliday GM (1987) Ventral tegmental (A10) system: neurobiology. 1. Anatomy and connectivity. Brain Res 434:117–165.
Paladini CA, Celada P, Tepper JM (1999) Striatal, pallidal, and pars reticulata evoked inhibition of nigrostriatal dopaminergic neurons is mediated by GABA(A) receptors in vivo. Neuroscience 89:799–812.
Phillips TJ, Huson M, Gwiazdon C, Burkhart-Kasch S, Shen EH (1995) Effects of acute and repeated ethanol exposures on the locomotor activity of BXD recombinant inbred mice. Alcohol Clin Exp Res 19:269–278.
Phillips TJ, Brown KJ, Burkhart-Kasch S, Wenger CD, Kelly MA, Rubinstein M, Grandy DK, Low MJ (1998) Alcohol preference and sensitivity are markedly reduced in mice lacking dopamine D2 receptors. Nat Neurosci 1:610–615.
Poncer JC, McKinney RA, Gahwiler BH, Thompson SM (2000) Differential control of GABA release at synapses from distinct interneurons in rat hippocampus. J Physiol (Lond) 528:123–130.
Rassnick S, Stinus L, Koob GF (1993a) The effects of 6-hydroxydopamine lesions of the nucleus accumbens and the mesolimbic dopamine system on oral self-administration of ethanol in the rat. Brain Res 623:16–24.
Rassnick S, D'Amico E, Riley E, Koob GF (1993b) GABA antagonists and benzodiazepine partial inverse agonists reduce motivated responding for ethanol. Alcohol Clin Exp Res 7:124–130.
Risinger FO, Cunningham CL (1995) Genetic differences in ethanol-induced conditioned taste aversion after ethanol pre-exposure. Alcohol 12:535–539.
Rozov A, Burnashev N, Sakmann B, Neher E (2001) Transmitter release modulation by intracellular Ca2+ buffers in facilitating and depressing nerve terminals of pyramidal cells in layer 2/3 of the rat neocortex indicates a target cell-specific difference in presynaptic calcium dynamics. J Physiol (Lond) 531:807–826.
Salin PA, Malenka RC, Nicoll RA (1996) Cyclic AMP mediates a presynaptic form of LTP at cerebellar parallel fiber synapses. Neuron 16:797–803.
Samson HH, Tolliver GA, Pfeffer AO, Sadeghi KC (1987) Oral ethanol reinforcement and the effect of the partial inverse benzodiazepine agonist RO 15-4513. Pharmacol Biochem Behav 33:601–608.
Samson HH, Hodge CW, Tolliver GA, Haraguchi M (1993) Effect of dopamine agonists and antagonists on ethanol-reinforced behavior: the involvement of the nucleus accumbens. Brain Res Bull 30:133–141.
Seamon KB, Daly JW (1985) High-affinity binding of forskolin to rat brain membranes. Adv Cyclic Nucleotide Protein Phosphor Res 19:125–135.
Self DW, Nestler EJ (1995) Molecular mechanisms of drug reinforcement and addiction. Annu Rev Neurosci 18:463–495.
Self DW, Genova LM, Hope BT, Barnhart WJ, Spencer JJ, Nestler EJ (1998) Involvement of cAMP-dependent protein kinase in the nucleus accumbens of cocaine self-administration and relapse of cocaine-seeking behavior. J Neurosci 18:1848–1859.
Smith BR, Robidoux J, Amit Z (1992) GABAergic involvement in the acquisition of voluntary ethanol intake in laboratory rats. Alcohol Alcohol 27:227–231.
Spanagel R, Weiss F (1999) The dopamine hypothesis of reward: past and current status. Trends Neurosci 22:521–527.
Stanford IM, Cooper AJ (1999) Presynaptic μ and δ opioid receptor modulation of GABAA IPSCs in the rat globus pallidus *in vitro*. J Neurosci 19:4796–4803.
Steffensen SC, Henriksen SJ, Wilson MC (1999) Transgenic rescue of SNAP-25 restores dopamine-modulated synaptic transmission in the coloboma mutant. Brain Res 847:186–195.
Steffensen SC, Nie Z, Criado JR, Siggins GR (2000) Ethanol inhibition of *N*-methyl-D-aspartate responses involves presynaptic gamma-aminobutyric acid (B) receptors. J Pharmacol Exp Ther 294:637–647.
Stoop R, Poo MM (1996) Synaptic modulation by neurotrophic factors: differential and synergistic effects of brain-derived neurotrophic factor and ciliary neurotrophic factor. J Neurosci 16:3256–3264.
Stuart GI, Redman SJ (1991) Mechanisms of presynaptic inhibition studied using paired-pulse facilitation. Neurosci Lett 126:179–183.
Sullivan JM (1999) Mechanisms of cannabinoid-receptor-mediated inhibition of synaptic transmission in cultured hippocampal pyramidal neurons. J Neurophysiol 82:1286–1294.
Terwilliger RZ, Beitner-Johnson D, Sevarino KA, Crain SM, Nestler EJ (1991) A general role for adaptations in G-proteins and the cyclic AMP system in mediating the chronic actions of morphine and cocaine on neuronal function. Brain Res 548:100–110.
Thiele TE, Willias B, Stadler J, Reynolds JG, Bernestein IL, McKnight GS (2000) High ethanol consumption and low sensitivity to ethanol-induced sedation in protein kinase A-mutant mice. J Neurosci 20:RC75:1–6.
Thomas MJ, Malenka RC, Bonci A (2000) Modulation of long-term depression by dopamine in the mesolimbic system. J Neurosci 20:5581–5586.
Waldeck RF, Pereda A, Faber DS (2000) Properties and plasticity of paired-pulse depression at a central synapse. J Neurosci 20:5312–5320.
Wan FJ, Berton F, Madamba SG, Francesconi W, Siggins GR (1996) Low ethanol concentrations enhance GABAergic inhibitory postsynaptic potentials in hippocampal pyramidal neurons only after block of GABAB receptors. Proc Natl Acad Sci USA 93:5049–5054.
Wand G, Levine M, Zweifel L, Schwindinger W, Abel T (2001) The cAMP-protein kinase A signal transduction pathway modulates ethanol consumption and sedative effects of ethanol. J Neurosci 21:5297–5303.
Weiss F, Parson LH, Schulteis G, Hyytia P, Lorang MT, Bloom FE, Koob GF (1996) Ethanol self-administration restores withdrawal-associated deficiencies in accumbal dopamine and 5-hydroxytryptamine release in dependent rats. J Neurosci 16:3474–3485.
Yoshimura M, Tabakoff B (1999) Ethanol's actions on cAMP-mediated signaling in cells transfected with type VII adenylyl cyclase. Alcohol Clin Exp Res 23:1457–1461.
Yun SH, Cheong MY, Mook-Jung I, Huh K, Lee C, Jung MW (2000) Cholinergic modulation of synaptic transmission and plasticity in entorhinal cortex and hippocampus of the rat. Neuroscience 97:671–676.
Zucker RS (1989) Short-term synaptic plasticity. Annu Rev Neurosci 12:13–31.